EXPANDED SITE INSPECTION REPORT

FOR

JACKPILE-PAGUATE URANIUM MINE PAGUATE, CIBOLA COUNTY, NEW MEXICO

Prepared for

U.S. ENVIRONMENTAL PROTECTION AGENCY

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EXECUTIVE SUMMARY

Weston Solutions, Inc. (WESTON®), the EPA Region 6 Superfund Technical Assessment and Response Team (START-3) contractor, has prepared this Expanded Site Inspection (ESI) Report to provide the U.S. Environmental Protection Agency (EPA) Region 6 with the background information collected for the site; discuss the ESI sampling activities; and present the analytical data obtained as part of the investigation performed at the Jackpile-Paguate Uranium Mine (Jackpile-Paguate) Site in Paguate, Cibola County, New Mexico.

The Site was identified as a potential hazardous waste site and entered into the Comprehensive Environmental Response, Compensation, and Liability Information System (CERCLIS) on 6 October 2009 (NMN000607033). A Preliminary Assessment (PA) was completed for the site on 29 November 2010, the results of which indicated that further sampling at the Site was warranted. START-3 conducted a Site Inspection (SI) in June 2010 that included the collection of soil, sediment, and surface water samples from the Site, including background and downgradient locations. These samples were analyzed for isotopic and total uranium, metals including mercury, and cyanide. Based on the analytical results, releases to surface water were documented.

Based on results of SI report, the EPA tasked an Expanded Site Assessment (ESI) of the Site to evaluate the Site using the ground water to surface water pathway. START-3 prepared a draft Conceptual Site Model (CSM) in January 2011 based primarily on the data and conclusions of the SI and other historical Site reports. The purpose of the CSM was to compile the existing hydrologic and geochemical data for the Site and identify data gaps that could be fulfilled within the budget and scope of an ESI. START-3 executed the ESI in April 2011 under the EPA-approved Jackpile-Paguate Expanded Site Inspection (ESI) Quality Assurance Sampling Plan.

Based on the 2011 ESI surface water and sediment sampling results, a release of isotopic uranium, isotopic thorium, gross alpha and beta, Radium 226 and 228, antimony, arsenic, beryllium, cadmium, cobalt, lead, nickel, and silver to the surface water pathway is documented. These contaminants were found at concentrations greater than significantly above those found in background samples collected.

This report	was prepared as part of the requirements of the Technical Direction Document (TDD)
No. TO-001	19-10-11-01 and serves as documentation of work completed to date.
	The EPA Site Assessment Manager did not provide final approval of this report prior to the completion date of the work assignment. Therefore, Weston Solutions, Inc. has submitted this report absent the Site Assessment Manager's approval.
X	The EPA Site Assessment Manager has provided final approval of this report. Therefore, Weston Solutions, Inc. has submitted this report with the Site Assessment Manager's approval.

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LIST OF ACRONYMS

BIA Bureau of Indian Affairs
BLM Bureau of Land Management

CERCLA Comprehensive Environmental Response, Compensation, and Liability Act CERCLIS Comprehensive Environmental Response, Compensation, and Liability

Information System

CoC contaminant of concern CSM Conceptual Site Model

EIS Environmental Impact Statement

EPA United States Environmental Protection Agency

ESI Expanded Site Inspection

GIS Geographical Information System

gpm gallons per minute
HRS Hazard Ranking System
mCi/l milliCuries per liter

MCL Maximum Contaminant Level mg/kg milligrams per kilogram mg/l milligrams per liter micrograms per liter

MSSL 2009 EPA Region 6 Human Health Medium Specific Screening Levels

NELAP National Environmental Laboratory Accreditation Program

NPL National Priority List
PA Preliminary Assessment
pCi/kg picocuries per kilogram
pCi/L picocuries per liter
PPE Probable Point of Entry

QASP Quality Assurance Sampling Plan

RCRIS Resource Conservation and Recovery Information System

ROD Record of Decision RSL regional screening level

SARA Superfund Amendments and Reauthorization Act of 1986

SI Site Inspection

START-3 EPA Region 6 Superfund Technical Assessment and Response Team

TAL Target Analyte List

TDD Technical Direction Document

TDL target distance limit
TDS total dissolved solids

USGS United States Geological Survey

WESTON Weston Solutions, Inc.
WHPA Well Head Protection Area

1. INTRODUCTION

1.1 BACKGROUND

Under the authority of the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA) and the Superfund Amendments and Reauthorization Act of 1986 (SARA), Weston Solutions, Inc. (WESTON®), the Environmental Protection Agency (EPA) Region 6 Superfund Technical Assessment and Response Team (START-3) contractor, has been tasked to perform an Expanded Site Inspection (ESI) of the Jackpile-Paguate Uranium Mine site ("the Site") located in Paguate, Cibola County, New Mexico (Figure 1-1). EPA Region 6 retained WESTON to complete this investigation under Contract Number EP-W-06-042, Technical Direction Document (TDD) No. TO-0019-10-11-01 (Appendix H).

The Site was identified as a potential hazardous waste site and entered into the Comprehensive Environmental Response, Compensation, and Liability Information System (CERCLIS) on 6 October 2009 (NMN000607033). A Preliminary Assessment (PA) was completed for the Site on 29 November 2010, the results of which indicated that further sampling at the Site was warranted. WESTON conducted a Site Inspection (SI) in June 2010 that included the collection of soil, sediment, and surface water samples from the Site, including background and downgradient locations. These samples were analyzed for isotopic and total uranium, metals including mercury, and cyanide. Based on the analytical results from the SI, an observed release to surface water was documented (Reference 24, Reference 25).

START-3 prepared a draft Conceptual Site Model (CSM) in January 2011 based primarily on the data and conclusions of the SI and other historical Site reports (Reference 27). The purpose of the CSM was to compile the existing hydrologic and geochemical data for the Site and identify data gaps that could be filled within the budget and scope of an ESI. START-3 executed the ESI in April 2011 under the EPA-approved Jackpile-Paguate ESI Quality Assurance Sampling Plan (Reference 8).

This document represents the draft report for the ESI. The purpose of this report is to provide the background information collected for the Site, discuss the ESI sampling activities, and present the analytical data obtained as part of the investigation.

1.2 OBJECTIVES OF THE INVESTIGATION

After reviewing the SI report, the EPA determined that an ESI would be necessary to help prioritize further work for the Site. The ESI is intended to be part of an on-going screening process of known and potential hazardous waste sites. Expanding on the information documented during the SI, the purpose of this ESI is to further identify immediate or potential threats that hazardous substances attributable to the Site may pose to human health and the environment. This is done by documenting the existence and migration of hazardous substances related to the Site and by identifying the receptors, or targets, potentially exposed to the hazardous substances. EPA will use the information obtained during the ESI to evaluate the Site using the Hazard Ranking System (HRS) and to help determine whether the Site is a potential candidate for inclusion on the National Priorities List (NPL). The intent of the ESI is to provide the documentation necessary to either rank the Site on the NPL or assign a "No Further Remedial Action Planned" status to the Site.

1.3 SCOPE OF WORK

The ESI Scope of Work is intended to describe the tasks requiring completion in order to evaluate the site using the HRS (Reference 1). As part of this ESI, START-3 performed the following major tasks:

- Prepared a site-specific Quality Assurance Sampling Plan (QASP), approved by the EPA, and Health and Safety Plan (HASP) prior to sampling activities.
- Gathered information documenting the environmental setting of the areas of concern to characterize the ground water, surface water, soil exposure, and air pathways.
- Reviewed available regulatory compliance files from federal, state, and local government agencies.
- Prepared a CSM to compile the existing hydrologic and geochemical data and study the ground water to surface water migration for the Site and identify data gaps indentified in the SI.
- Conducted ESI field sampling activities on 19 April 2011 through 21 April 2011. Samples were collected within ground water and surface water locations at the Site and within potential pathways of migration. The samples were collected in general accordance with the site-specific QASP and HASP to document the presence and migration of hazardous substances attributable to the site.

- Evaluated the available information from the on-site observations, historical aerial photographs, property ownership records, interviews, area environmental information, and area demographic characteristics.
- Submitted the ESI samples to National Environmental Laboratory Accreditation Program (NELAP) certified laboratories for analysis and reviewed and tabulated the resulting data.
- Compared the laboratory results to 2009 EPA Region 6 Human Health Medium-Specific Screening Levels (MSSLs), EPA Maximum Contaminant Levels (MCLs), and three times the background concentrations (Appendix F, Reference 1).
- Prepared this report to present the findings of the ESI.

1.4 REPORT FORMAT

The ESI report is presented in a format that is intended to facilitate evaluation of the site using the HRS (Reference 1). The report contains the following sections:

- Section 1 Introduction
- Section 2 Site Characteristics
- Section 3 Investigative Efforts
- Section 4 Waste Source Characterization
- Section 5 Ground Water Pathway
- Section 6 Surface Water Pathway
- Section 7 Soil Exposure and Air Migration Pathways
- Section 8 Summary
- Section 9 References

Additional information is provided in the appendices following the text of the report. The appendices are as follows:

- Appendix A Digital Photographs
- Appendix B START-3 Site Logbooks
- Appendix C START-3 Quality Assurance Sampling Plan
- Appendix D Laboratory Data Packages
- Appendix E Laboratory Data Validation Packages
- Appendix F EPA National Primary Drinking Water Regulations (MCLs) and 2009 EPA Region 6 Human Health Medium-Specific Screening Levels (MSSLs)
- Appendix G Reference Documentation

THIS DOCUMENT WAS PREPARED BY WESTON SOLUTIONS, INC., EXPRESSLY FOR EPA. IT SHALL NOT BE RELEASED OR DISCLOSED IN WHOLE OR IN PART WITHOUT THE EXPRESS, WRITTEN PERMISSION OF EPA.

• Appendix H TDD No. 0019-10-11-01 and Amendments A-C

The tables referred to throughout sections of this report are provided within the sections in which they are first cited or as separate portable document format (PDF) files. Figures cited in this report are provided as PDF files. Photographs taken during the ESI field sampling activities are provided as Appendix A. The START-3 field logbook notes are provided as Appendix B.

2. SITE CHARACTERISTICS

2.1 LOCATION

The Site is a former uranium mine located on the Pueblo of Laguna, approximately 40 miles west of Albuquerque in Paguate, Cibola County, New Mexico (Reference 2, Reference 3, Reference 23, Reference 30). The Site is located at Latitude 35° 08' 16.74" North and Longitude 107° 20' 51.84" West. Latitude and longitude coordinates were measured from the approximate center of the former mine and were determined using a scaled aerial photograph and geographical information system (GIS) Software. The Site is located in an area of canyons and arroyos to the east of the village of Paguate and can be reached by taking SR 124 off of Interstate 40, 1.7 miles west, then turning right or north on SR 279 and continuing approximately 6.5 miles. The Site location is shown in Figure 1-1.

The town of Paguate is located adjacent and west of the Site with a population of approximately 492. St. Anthony Mine (the closest uranium mining operation) is located approximately 11,000 feet northeast of the Site (Reference 2, Reference 4, Reference 23, Reference 30).

2.2 SITE DESCRIPTION

The property on which the former uranium mine is located encompasses approximately 7,868 acres. Approximately 2,656 acres of this property were disturbed by mining operations including three open pits (South Paguate, North Paguate, and Jackpile) that were between 200 and 300 feet deep, 32 waste dumps, and 23 protore (rock and soil associated with ore body) stockpiles (Reference 3, Reference 5). The Site layout is shown in Figure 2-1.

The remaining surrounding area is mainly undeveloped and consists of a canyon/mesa topography containing wildlife consisting of elk, antelope, goats, and mule deer; in addition, the Pueblo of Laguna utilizes the surrounding lands for cattle grazing and watering. The perennial rivers Rio Moquino and Rio Paguate bisect the Site near its center. The Rio Moquino flows southeasterly into the Rio Paguate within the Site, and the Rio Paguate continues to flow south of the Site through the Paguate Reservoir and joins the Rio San Jose approximately 1 mile southwest of the reservoir (Figure 4-2, Reference 2, Reference 23, Reference 30).

2.3 OPERATIONAL HISTORY

The Site is part of the Grants Uranium Mineralization Belt that covers much of northwestern New Mexico. The Jackpile-Paguate Mine operated from 1953 to 1982, primarily as an open-pit mine. Subsurface mining also occurred in the southwestern corner of the property from 1974 to 1980, extending from the South Paguate Pit southward under old Highway 279. The mine was operated by the Anaconda Mineral Company (later Atlantic Richfield Company). During the course of mining activities, 356 million tons of material was disturbed, and 22 million tons of ore were removed from the Site. The mine closed in 1982 due to the fall in the price of uranium (Reference 3, Reference 5, Reference 28, Reference 29).

In December 1986, under a series of agreements between the Bureau of Indian Affairs (BIA) and the Pueblo of Laguna, it was agreed that the Pueblo of Laguna would perform management, coordination, and administration of the Jackpile-Paguate Reclamation Project in accordance with the requirements set forth in the Jackpile-Paguate Environmental Impact Statement (EIS) and the associated Record of Decision (ROD) (Reference 3, Reference 5).

Reclamation of the Jackpile-Paguate Uranium Mine commenced in 1990. The "preferred alternative" reclamation plan incorporated the following: backfilling the open pit areas to at least 10 feet above projected ground water recovery levels using protore and waste rock dump material; reducing the upper 15 feet of highwall slopes; recontouring and covering remaining waste rock dumps; completing arroyo drainage improvements and erosion controls; decontaminating remaining structures; removing/disposing of non-essential structures; plugging and bulkheading underground ventilation raises and decline portals, respectively; reclaiming miscellaneous features such as wells, access roads, rail spurs, drill holes, etc.; performing site-wide revegetation of disturbed areas; and providing site security and long-term monitoring of reclamation success for a period of not less than 10 years (Reference 5).

On 10 June 1995 the Laguna Construction Company, which was in charge of the reclamation, officially closed the Jackpile Reclamation Project (Reference 7).

In September 2007, a ROD Compliance Assessment for Jackpile-Paguate Uranium Mine was completed by OA Systems Corporation to determine if the post-reclamation had met the

requirements of the EIS and ROD. This report concluded that reclamation of the mine was still not complete as several non-compliant and potentially non-compliant issues still needed to be addressed. The mine would be considered complete when revegetated sites reached 90 percent of the density, frequency, foliar cover, basal cover, and production of undisturbed reference areas (but no sooner than 10 years following seeding). In addition, gamma radiation levels were to be no greater than twice background over the entire mine site. Outdoor radon 222 concentrations were to be no greater than 3 picocuries per liter (pCi/L). Radon daughter levels in any remaining surface facilities were not to exceed 0.03 Working Levels (WL) (Reference 5).

2.4 REGULATORY INVOLVEMENT

The Site is not listed in the Resource Conservation and Recovery Information System (RCRIS) database, as of 11 July 2011 (Reference 26). The Site is listed in CERCLIS as NMN000607033 (Reference 25).

3. INVESTIGATIVE EFFORTS

3.1 PREVIOUS SAMPLING

3.1.1 Anaconda Remedial and Investigative Efforts

Anaconda conducted a hydrologic investigation at the Site during the years 1977 through 1979, while the mine was still operating. The purpose was to determine whether accumulated pond water from two specific holding ponds (Rabbit Ear Pond and P-10 Pond) posed a threat to ground water and/or surface water. To address these issues, eleven wells were installed across the active operational area of the Site, screened in the Alluvial Aquifer, the Jackpile Sandstone, and a number of waste piles. The hydrologic model of the Site provided a ground water flow model for the Site, which the U.S. Geological Survey (USGS) later used to construct its more detailed hydrologic model (Reference 9, Reference 29, Reference 31).

In 1982, Hydro Geo Chem, Inc. prepared a report for the Pueblo of Laguna that examined the effects of uranium mine dewatering on the local water resources. The report described dewatering simulations from the five regional mining operations, including the Jackpile Paguate pits. The results of the simulations indicated significant (20 to 200 feet) drawdown of the ground water surface near the mine pits due to dewatering activities at the mines. The report also stated that water in the lower Rio Paguate and Rio Moquino is thought to be contaminated by activities at the Jackpile mine (Reference 21).

The Anaconda Minerals Company conducted annual monitoring through 1986, as evidenced by Data Transmission files available at the Bureau of Land Management (BLM). These data include chemical and radiologic analyses for surface water and ground water across the Site, as well as air sampling and radon monitoring (Reference 18, Reference 19, Reference 20).

The Anaconda Mineral Company commissioned an evaluation of the hydrologic effects of the proposed backfilling of the North Paguate, South Paguate, and Jackpile pits. A 1983 report indicated that ground water levels in pits would recover to a steady state in 30, 150, and 300 years, respectively. The report recommended the installation of fine-grained materials along the upgradient margin to cut off ground water flow into the North Paguate Pit to keep the water levels

below the proposed level of the backfill. The report noted that it was important to keep water levels in the pits low in order to slow the oxidation process, as well as prevent water from ponding in the pits (Reference 6).

The 1983 report predicted that uranium and radium concentrations would initially increase to levels exceeding 5 milligrams per liter (mg/l) and/or 30 milliCuries per liter (mCi/l) as the backfill became saturated, but these levels would decrease as the oxygen levels were depleted. Ground water flow from the North Paguate Pit was predicted to flow to the southeast, discharging to the Alluvial Aquifer at a rate of 20 gallons per minute (gpm). Ground water flow from the Jackpile Pit was predicted to flow south/southwesterly, discharging to the Alluvial Aquifer at a rate of 1.5 gpm. Flow from the South Paguate Pit was expected to flow predominantly to the east/southeast through the Jackpile sandstone to where it outcrops in the cliffs above the Paguate River; a smaller quantity of South Paguate Pit waters are expected to flow to the Paguate River at a rate of 1.3 gpm. The report concluded that dissolved solids and radiological materials will have a negligible impact on surface water quality, based on these numbers (Reference 6).

The USGS prepared a Regional Water Resources report in 1983 that provided general surface water and ground water yield data for the Pueblo of Laguna. The report underscored the Pueblo of Laguna's reliance on surface water and ground water resources for agricultural irrigation, as well as livestock watering and public consumption (Reference 12).

The USGS conducted aquifer tests on paired wells installed on the Site. The well pairs included the steel-cased, M-series wells, and paired PVC-cased wells installed at the same hydrologic interval. Constant discharge tests were conducted on three sets of paired wells screened in the Jackpile Sandstone (M-2, M-3, and M-21) and one set of paired wells in the Alluvial Aquifer (M4). An additional slug test was performed at M-25, which is screened in a lower sand unit in the Brushy Basin Shale Member. Transmissivity values for the Jackpile Sandstone were observed in the range of 2.0 to 47 square feet per day; transmissivity for the Alluvial Aquifer was calculated at 430 square feet per day. Hydraulic conductivity for the Jackpile Sandstone was reported in the range of 0.025 to 0.39 feet per day; hydraulic conductivity for the Alluvial Aquifer was reported at 23 feet per day. The discharge test at M-4 also indicated that water levels in the underlying Jackpile Sandstone were affected by the test, indicating hydraulic communication between these units (Reference 11).

The USGS published "Hydrology and Water-Quality Monitoring Considerations, Jackpile Uranium Mine, Northwestern New Mexico," in 1985. The report stated that Rio Paguate was a gaining stream prior to mining activities, during the period of 1977-1980. However, Rio Paguate has been losing water at the confluence with Rio Moquino, likely due to depressed aquifer conditions at the Jackpile and Paguate pits. The report further stated that backfill of the pits would result in saturation, and possible oxidation by ground and surface water in the backfill may cause increased solution of rock minerals, including radionuclides and trace elements. Discharge from the backfill in the Paguate pits will flow through the waste rock piles to Rio Paguate. Discharge from the backfilled Jackpile pit would likely flow to the Jackpile Sandstone, then to the Alluvial Aquifer, then to the Rio Paguate. The report recommended additional monitoring locations in the downgradient areas for water quality parameters and radionuclides (Reference 29).

As part of a ROD Compliance Assessment, surface water data at the Site were evaluated for 10 years between 1997 and 2007. During this evaluation, several samples were found to exceed EPA Maximum Contaminant Levels (MCLs) for drinking water for gross alpha radiation particles (MCL -15 pCi/l) and total uranium (MCL - 0.03 mg/l) and the National Secondary Drinking Water Standard of 0.05 mg/l for manganese (Reference 3). The following concentration ranges were found in samples collected during the 10-year ROD compliance timeframe:

Table 3-1						
ROD	Compliance Assessme	ent Analytical Summar	y			
Location	Gross Alpha Radiation Concentrations pCi/l	Total Uranium Concentrations mg/l	Manganese Concentrations mg/l			
Rio Paguate – Ford Crossing-Rail Trestle	37.9 - 214.33	0.08 - 544.14	<0.01 - 10.7			
Lower Rio Moquino	5.24 - 53.05	0.03 - 234.95	0.0 - 0.127			
Lower Rio Paguate	2.24 - 106.22	0.016 - 149.62	0.0357 - 0.13			
Paguate-Moquino Confluence	[-21.98 ¹] - 94.03	0.029 - 252.59	0.0112 - 0.14			
Upper Rio Moquino	$[-12.46^{1}] - 35.11$	0.008 - 52.89	<0.01 - 0.036			
Upper Rio Paguate	ND - 25.53	0.002 - 32.21	0.0396 - 0.146			
Paguate Reservoir	ND - 3.04	0.002 - 17.42	<0.01 - 0.004			

Notes:

pCi-picocuries per liter mg/l – milligrams per liter

As part of the Water Pollution Program Grants (Section 106), the Pueblo of Laguna Environmental and Natural Resources Department collected surface water samples along the surface water pathway quarterly from 2005 through 2009. The samples were analyzed for isotopic uranium and total uranium. As with the samples collected as part of the ROD Compliance Assessment, these samples also contained total uranium concentrations exceeding the EPA MCL of 0.03 mg/l which is equivalent to 30 micrograms per liter (μ g/l) (Reference 14). The following uranium concentration ranges were found in the samples collected:

¹ Negative counts [] may occur since sample counts are compared to background counts, and background counts reflect naturally-occurring radionuclides and cosmic radiation that are detected by laboratory instrumentation. Samples that are not different from background may have a negative value when background is subtracted (Reference 13).

ND - Not Detected.

< - Less Than.

Table 3-2							
Water Polluti	Water Pollution Program Quarterly Sampling Result Summary						
Location	Total Uranium Concentrations µg/l	U-234 Concentrations μg/l	U-235 Concentrations µg/l	U-238 Concentrations µg/l			
RPG-03 Rio Paguate below the former Jackpile Mine (at the PPE)	0.007 - 425.557	0.004 - 92.667	$[-0.032^1] - 4.72$	0.003 - 448			
RPG-04 Rio Paguate at Mesita Dam (within the TDL)	0.008 - 36.533	0.008 - 72.939	[-0.008 ¹] - 2.208	0.039 - 216			
RSJ-04 Rio San Jose below the Dipping Vat Spring on Sedillo Grant (beyond the TDL)	2.012 - 21.615	0.001 - 9.435	0.043 - 0.547	0.675 - 14.6			
RMQ-01 Rio Moquino above Jackpile Mine (background)	0.005 - 31.138	0.000 - 5.972	0.000 - 1.177	0.005 - 30.904			

Notes:

PPE = Probable Point of Entry

3.1.2 EPA Preliminary Assessment (PA) and Site Investigation (SI)

START-3 conducted a PA at the Site in April 2010, the results of which indicated that further sampling at the Site was warranted. START-3 completed a SI in June 2010, which included sampling and chemical/radiological analyses of waste piles, sediment, soil, and surface water at the Site. The results of the SI indicated the presence of hazardous substances at the Site, as well as a release of hazardous substances from the Site to surface water (Reference 24).

START-3 initiated the SI field sampling activities on 2 March 2010. The field sampling activities were completed in general accordance with the EPA-approved site-specific Quality Assurance Sampling Plan (QASP) dated February 2010. START-3 followed START-3 Sample Operating Procedures for surface soil sampling, stockpile sampling, surface water sampling, and sediment sampling (Reference 8, Reference 24). Sampling locations and results are presented in Figure 3-1 and Tables 3-3 through 3-6.

¹ Negative counts [] may occur since sample counts are compared to background counts, and background counts reflect naturally occurring radionuclides and cosmic radiation that are detected by laboratory instrumentation. Samples that are not different from background may have a negative value when background is subtracted (Reference 13).

 $TDL = Target\ Distance\ Limit$

µg/l – microgram per liter

3.1.2.1 Source Sampling

In an effort to characterize the possible sources associated with the Site, START-3 collected four soil samples (includes one background soil sample), one sediment sample, and one surface water sample from locations in proximity to open pits, waste dumps, and protore stockpiles throughout the site. Sampling locations are presented in Figure 3-1. The samples were analyzed for isotopic and total uranium and Target Analyte List (TAL) metals with mercury and cyanide. The analytical results were compared to 2009 EPA Region 6 Human Health Medium Specific Screening Levels (MSSLs). The soil and sediment samples were compared to residential soil levels, and the water sample was compared to MCLs (Reference 24).

Laboratory analytical results for the six source samples collected from sample stations JM-01, JM-02, JM-03, and JM-05 were used to characterize the possible sources at the site. One soil sample, JM-SS-04, was collected at the northern edge of the mine to establish the soil background concentrations in the area. The background sample was collected to establish that contamination within the source soil samples was from Jackpile-Paguate mining operations and not from a release from outside sources. This location was chosen because it is upgradient and upstream of the Site and in close proximity to other mining operations in the area. St. Anthony Mine is located approximately 11,000 feet northeast of the Jackpile-Paguate Mine (Reference 2).

Total uranium, chromium, manganese, nickel, selenium, and vanadium were all present at concentrations significantly above background. The concentration of total uranium exceeded EPA MSSLs and MCLs in samples JM-SED-02, JM-SS-05, and JM-SW-02 (Reference 24). Sampling results are presented in Tables 3-3 and 3-4.

	Table 3-3 Summary of SI Source Soil/Sediment Analytical Data							
Analyte	Units	Residential Soil MSSL	JM-SS-04 Background	JM-SS-01	JM-SED-02	JM-SS-03	JM-SS-05	
U-232	mg/kg	ı	1.86E-07	1.64E-07	1.55E-07	6.07E-07	1.77E-06	
U-234	mg/kg	-	0.00066	0.00173	0.0226	0.0064	<u>0.0144</u>	
U-235	mg/kg	-	0.084	0.343	<u>2.55</u>	<u>0.94</u>	<u>2.06</u>	
U-238	mg/kg	1	12.5	<u>40.8</u>	<u>362</u>	<u>126</u>	<u>271</u>	
Uranium, Total	mg/kg	230	12.5	<u>41.1</u>	<u>364</u>	<u>127</u>	<u>273</u>	
Aluminum	mg/kg	77000	2400	1900	6700	3100	1900	
Arsenic	mg/kg	0.39	2.9	2.9	<u>10</u>	2.9	1.6	
Barium	mg/kg	15000	22	53	32	58	30	
Calcium	mg/kg	1	5000	930	<u>40000</u>	8700	910	
Chromium	mg/kg	1	1.6	<1	5.8	3.2	1.2	
Cobalt	mg/kg	23	3.2	1.7	5.7	2.9	<1	
Copper	mg/kg	3100	9.5	3.2	8.2	3.9	1.2	
Iron	mg/kg	55000	6300	3400	17000	9000	3700	
Lead	mg/kg	400	6.4	5.4	10	8.3	9.2	
Magnesium	mg/kg	-	2900	780	<u>13000</u>	3000	730	
Manganese	mg/kg	1800	50	16	<u>270</u>	84	25	
Nickel	mg/kg	-	3.5	<2	<u>11</u>	4.5	<2	
Potassium	mg/kg	ı	470	350	<u>1700</u>	630	370	
Selenium	mg/kg	390	<0.51U	<u>1.6</u>	3.4	<u>1.9</u>	< 0.51	
Sodium	mg/kg	-	<100U	<100	2200	<u>270</u>	<100	
Vanadium	mg/kg	390	3.7	<u>30</u>	<u>15</u>	20	<u>90</u>	
Zinc	mg/kg	23000	46	4.9	36	16	3.3	

Notes:

 $MSSL:\ 2009\ EPA\ Region\ 6\ Human\ Health\ Medium\ Specific\ Screening\ Levels.$

mg/kg: milligrams per kilogram.

<: analyte not present at concentration above the associated reporting limit.

Bold and underlined values are significantly above the background concentration.

Highlighted values are significantly above background and exceed MSSLs.

Table 3-4							
Summary of SI Source Surface Water Analytical Data							
Analyte	Units	MCL	JM-SW-02				
U-232	μg/l	-	0.0000084				
U-234	μg/l	-	0.538				
U-235	μg/l	-	61				
U-238	μg/l	-	8500				
Uranium, Total	μg/l	30	8500				
Aluminum	μg/l	-	3500 JL				
Calcium	μg/l	-	750000 JL				
Cobalt	μg/l	-	23 JL				
Iron	μg/l	-	10000 JL				
Magnesium	μg/l	-	1900000 JL				
Manganese	μg/l	-	1300 JL				
Potassium	μg/l	-	57000 JL				
Sodium	μg/l	-	1400000 JL				
Zinc	μg/l	-	26 JL				

Notes:

MCL: Maximum Contaminant Level. JL: Estimated Concentration Biased Low. µg/l: micrograms analyte per liter water. Highlighted values exceed MCLs.

3.1.2.2 Surface Water/Sediment Sampling

START-3 collected 15 co-located surface water and sediment samples (total of 30 samples), including 3 co-located surface water and sediment background samples (RM- BG, RP- BG, RSJ-BG), and 2 surface water field duplicate samples (RP-SW-BGD and PR-SE-01D), and 2 sediment field duplicate samples (RP-SED-BGD and PR-SED-01D). These samples were collected in an effort to characterize the surface water pathway and to document an observed release to surface water attributable to the Site. Sampling locations are presented in Figure 3-1. The surface water and sediment samples were analyzed for isotopic uranium, total uranium, and TAL metals with mercury and cyanide. The analytical results were compared to 2009 EPA Region 6 MSSLs. The sediment samples were compared to residential soil levels and the water samples were compared to the MCLs (Reference 24).

None of the sediment samples collected within the surface water pathway contained results significantly above the background concentrations or above the MSSLs. Several surface water

sample results significantly exceeded background and MCL concentrations. Samples RP-JM-SW-01, RP-SW-01D, RP-SW-02, and PR-SW-01 contained uranium isotopes U-234, U-235, U-238 and total uranium concentrations significantly greater than the background concentrations. Samples RP-SW-01, RP-SW-01D, and PR-SW-01 contained manganese concentrations significantly above background concentrations. Samples RP-SW-01, RP-SW-01D, RP-SW-02, RP-SW-03, and PR-SW-01 contained total uranium concentrations above MCLs (Reference 24). Sampling results are presented in Tables 3-5 and 3-6 for sediment and surface water, respectively.

3.2 EXPANDED SITE INSPECTION SAMPLING

START-3 was tasked to conduct an ESI of the Site, including the collection of environmental samples. The specific sampling objectives were to collect data that could be used to document a release of hazardous substances to the surface water or ground water pathway. The site-specific Contaminants of Concern (CoCs) include isotopic uranium, total metals (including mercury), gross alpha, beta, and gamma, and radium 226 + 228. In addition, water samples were analyzed for total dissolved solids and general cation/anion chemistry. These data will be used to evaluate sample locations with the CSM, submitted under separate cover, that considers hydrologic and hydrochemical data. The field work was conducted in accordance with the START-3 QASP, approved by the EPA in April 2010 (Appendix C).

START-3 implemented the QASP during 19-21 April 2011. START-3 collected ground water samples from existing monitoring wells around the Site. Additionally, surface water and sediment samples were collected from the Rio Moquino, Rio Paguate, and the Paguate Reservoir. Background ground water, surface water, and sediment samples were collected from locations upgradient from mining activities. To support the ESI, samples were collected from 12 ground water locations, 9 sediment locations, and 8 surface water locations. Sample locations are listed in Table 3-7 and are presented in Figure 3-2.

Surface water, ground water, and surface sediment samples were collected and submitted to a National Environmental Laboratory Accreditation Program (NELAP) certified laboratory for the following analyses: isotopic uranium (including isotopes 234, 235, and 238) by ASTM 3972-90M alpha spectrometry; total metals including mercury by SW846 Methods 6010B and 7470/7471; gross alpha, beta, and gamma radiation by EPA Method 9310; and radium 226 +228 by EPA

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Method 903.1. In addition, water samples were analyzed for total dissolved solids (TDS) by EPA Method 160.1 and General Anion Chemistry by EPA Method 9056. The analytical data were validated by START-3. Data are presented in Tables 3-8 and 3-9. The validated laboratory data packages are included in Appendix E.

	Table 3-7 ESI Sample Locations and Sampling Rationale						
Sample Name	Sample Matrix	Sample Location (refer to Figure 3-2)	Rationale				
RM-SW-BG RM-SED- BG	Surface water Sediment	Rio Moquino upstream of Jackpile Mine.	Previous sample location. Collected to establish background concentration on Rio Moquino.				
RP-SW-BG RP-SED-BG	Surface water Sediment	Rio Paguate upstream of Jackpile Mine.	Previous sample location. Collected to establish background concentration on Rio Paguate.				
RM-JM-SW RM-JM- SED	Surface water Sediment	Rio Moquino within the mine boundaries, upstream of its confluence with Rio Paguate.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water.				
RP-JM-SW RP-JM-SED	Surface water Sediment	Rio Paguate within the mine boundaries, upstream of its confluence with Rio Moquino.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water.				
RP-JM-SW- 01 RP-JM- SED-01	Surface water Sediment	Rio Paguate after the confluence of it with Rio Moquino and within mine boundaries.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water.				
RP-SW-01 RP-SED-01	Surface water Sediment	Rio Paguate on the southern end of the mine boundary.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water.				
RP-SW-01 RP-SED-01	Surface water Sediment	Rio Paguate on the southern end of the mine boundary.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water.				
RP-SW-02* RP-SED-02	Surface water Sediment	Rio Paguate downstream of Jackpile Mine.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water.				
RP-SW-03* RP-SED-03	Surface water Sediment	Rio Paguate downstream of Jackpile Mine.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water.				

Table 3-7 ESI Sample Locations and Sampling Rationale (Continued)						
Sample Name	Sample Matrix	Sample Location (refer to Figure 3-2)	Rationale			
PR-SW-01 PR-SED- 01	Surface water Sediment	Paguate Reservoir (aka Mesita Reservoir) downstream of Jackpile Mine.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water.			
MD-SW (sample not part of ESI and data not included in Tables 3-8 & 3- 9)	Surface water	Irrigation canal off the Mesita Reservoir (aka Paguate Reservoir) downstream of Jackpile Mine. (not on figure – not part of ESI sampling).	This sample location is being collected per request from the Village of Mesita and results not part of ESI.			
MW-RM	Ground water	Monitoring well located off Rio Moquino, upgradient of Jackpile Mine.	Collected to establish background concentration upgradient and north of Jackpile Mine.			
MW-1	Ground water	Monitoring well located upgradient and to the northwest of Jackpile Mine.	Collected to establish background concentration upgradient of Jackpile Mine.			
MW-8	Ground Monitoring well located		Collected to establish background concentration upgradient of Jackpile Mine.			
SPOP35	Ground water	Monitoring well located within South Paguate Pit.	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations.			
Well#4	Ground water	Monitoring well located within Jackpile Mine along Rio Paguate.	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations.			
MW-3	Ground water	Monitoring well located within Jackpile Mine along Rio Paguate.	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations.			
NPOP20E	Ground water	Monitoring well located within North Paguate Pit.	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations.			
JPOP41N	Ground water	Monitoring well located within Jackpile Pit.	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations.			

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	Table 3-7 ESI Sample Locations and Sampling Rationale (Continued)							
Sample Name			Rationale					
MW-2	Ground water	Monitoring well located just south and downgradient of mine.	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in ground water with respect to background concentrations.					
MW-6	Ground water	Monitoring well located south and downgradient of mine.	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in ground water with respect to background concentrations.					
MW-MD	Ground water	Monitoring well located downgradient of mine and west of Mesita Dam.	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in ground water with respect to background concentrations.					

Notes:

3.2.1 Ground Water Pathway Sampling

Fourteen ground water samples (including two duplicates) were collected from existing monitoring wells. The ground water samples were submitted to a NELAP certified laboratory, for the following analyses: isotopic uranium (including isotopes 234, 235, and 238) by ASTM 3972-90M alpha spectrometry; total metals including mercury by SW846 Methods 6010B and 7470/7471; Gross alpha, beta, and gamma radiation by EPA Method 9310; Radium 226 and 228 by EPA Method 903.1; TDS by EPA Method 160.1; and General Anion Chemistry by EPA Method 9056 (Appendix D). Sampling locations are presented in Table 3-7 and Figure 3-2. Sample results are presented in Tables 3-8 and 3-9.

Isotopic uranium, isotopic thorium, gross alpha and beta radiation, Radium 226 and 228, beryllium, cadmium, chromium, cobalt, lead, nickel, selenium, silver, and zinc were all present in ground water at concentrations significantly above background.

3.2.2 Surface Water Pathway Sampling

START-3 collected 8 surface water and 10 sediment samples (including two background and one duplicate sample of each matrix) as part of the ESI to document an observed release to the surface water pathway attributable to the Site. In addition, one surface water sample was collected at station MD-SW, but this sample collection was not part of the ESI sampling event and data for this sample

^{* -} Sample locations were dry; therefore no surface water samples were collected from these locations.

is not discussed in this report. The surface water and/or sediment sample locations are presented in Table 3-7 and Figure 3-2. Analytical results are presented in Tables 3-8 and 3-9.

The sediments collected during the ESI were precipitates, rather than clastic sediments, in order to determine whether uranium might be transported and deposited as a precipitate, rather than as physically-eroded materials.

Isotopic uranium, gross alpha and beta radiation, arsenic, beryllium, cobalt, lead, nickel, and silver were all present in surface water at concentrations significantly above background. Isotopic uranium, isotopic thorium, gross alpha and beta radiation, Radium 226 and 228, antimony, and cadmium were all present in sediments and/or surface waters at concentrations significantly above background.

3.2.3 Deviations from the QASP

The following deviations from the QASP occurred during the field work:

- Isotopic Thorium by ASTM 3972-90M alpha spectrometry was added to the analyte list. This was done at the suggestion of BLM and the USGS in order to assist with future fate-and-transport studies.
- Nitrates were added to the analyte list for General Anion Chemistry by EPA Method 9056. This analyte was added at the request of the USGS in order to model oxidationreduction mixing in the surface water and ground water reservoirs.

None of these deviations are expected to have any adverse impact on the data or conclusions of this report.

4. WASTE SOURCE CHARACTERIZATION

Potential hazardous substance sources associated with the Site include, but may not be limited to:

- Open pits, waste dumps, and protore stockpiles associated with the Site. Mining operations at the Site resulted in 2,656 acres of surface disturbance, of which the open pits accounted for 1,015 acres. The Jackpile, North Paguate, and South Paguate open pits make up approximately 40 percent of the total disturbed acreage at the Site. Located in and around the open pits were 23 protore stockpiles. The protore stockpiles accounted for 103 acres of the disturbed area. The mine contained 32 waste dumps that accounted for approximately 48 percent of the disturbed area. Estimated quantities of wastes from the sources have been documented and include 1,105 acres of material in the open pits, 1,266 acres of material in the waste dumps, and 10.3 million cubic yards of material in the protore stockpiles (Reference 3).
- Approximately 23 million tons of uranium resources remain at the mine site as stockpiled protore and unmined deposits. Protore is material that was stockpiled throughout the mining operation because it contains elevated but sub-economic uranium concentrations. Approximately 21 million tons of protore containing 0.02 to 0.059 percent uranium (U-308) exist at the mine site. This material is located on the surface in 23 stockpiles dispersed throughout the mine. Approximately 2 million tons of unmined deposits containing 0.094 to 0.30 percent uranium (U308) remain at the Site. These resources are located in 11 deposits, three of which contain 90 percent of the resources (Reference 3).
- Wastes generated at the Jackpile-Paguate Uranium Mine were not removed from the Site but used for backfill. There was no impermeable cap placed on top of the backfill other than 3 feet of overburden (material removed during surface mining and stockpiled) and 2 feet of topsoil (Reference 3, Reference 5).
- Uranium was detected in Site sources at concentrations significantly above background, up to 364 milligrams per kilogram (mg/kg), during the March 2010 SI sampling. Sources also contained chromium, magnesium, manganese, nickel, selenium, and vanadium at concentrations significantly above background (Reference 24).

5. GROUND WATER PATHWAY

A discussion of the ground water pathway, one of four major pathways of potential hazardous waste migration assessed in this report, is provided in this section. The discussion focuses on the aquifer characteristics of the region, the likelihood of release to the ground water pathway, and the potential targets of the waste migration through the ground water pathway.

5.1 HYDROLOGIC DESCRIPTION

The Jackpile-Paguate Site is located in mesa and canyon country typical of much of the southeastern Colorado Plateau. It is situated in a broad valley of northwest-dipping, sandstone-capped benches pierced by numerous basaltic volcanic necks that rise up to 1,000 feet above the surrounding terrain. Sedimentary rocks exposed in the area of the Site range in age from late Triassic to Late Cretaceous. In addition, Tertiary-age diabase dikes and sills and volcanic flow rocks are exposed near the Site. At the Site, all of the rock units above the lower Mancos Shale have eroded. The stratigraphy of the mine includes the Morrison Formation, Dakota Sandstone, Mancos Shale, Tertiary igneous dikes, and Quaternary alluvium (Reference 3).

5.2 GROUND WATER CONDITIONS

According to the Water Well Report provided by GeoSearch on 14 April 2010, the nearest domestic well is located approximately 2.5 miles north of the Jackpile-Paguate Uranium Mine Site (Figure 4-1, Reference 10). GeoSearch identified a total of 11 ground water wells within a 4-mile radius of the Site. Water from these wells was used for the following purposes: domestic use (2 wells), monitoring wells (3 wells), water for mining, milling, or oil (5 wells), and sanitary in conjunction with commercial use (1 well). GeoSearch conducted the water well search using the United States Geological National Water Information System, and the Water Administration Technical Engineering Resource System (Reference 10). No Wellhead Protection Area (WHPA) has been identified within 4 miles of the Site. Ground water from the Alluvium and Jackpile Sandstone Aquifers is used for irrigation and livestock purposes (Reference 3).

5.3 LIKELIHOOD OF RELEASE

A release to ground water attributable to the Site is documented based on the results of the ESI. Isotopic uranium, isotopic thorium, gross alpha and beta radiation, Radium 226 and 228, beryllium, cadmium, chromium, cobalt, lead, nickel, selenium, silver, and zinc were all present in ground water at concentrations significantly above background.

5.4 GROUND WATER PATHWAY CONCLUSIONS

Although a release has been determined, the mine floor is generally above the water table and there are no wells located downgradient (south) of the site (Reference 22).

6. SURFACE WATER PATHWAY

Surface water is the second of four pathways of potential hazardous waste migration assessed for the Site. A discussion of the types of surface water draining the Site, the Probable Point of Entry (PPE) for a hazardous substance from the Site to enter the surface water, the likelihood of a release, and the potential targets of the pathway are discussed in this section.

6.1 HYDROLOGICAL SETTING

There is no overland flow segment from the source to the in-water segments. The Rio Moquino and Rio Paguate bisect the mine and are in direct contact with Site sources (Reference 2, Reference 9). Within the mine footprint, ground water in the Jackpile Sandstone Aquifer interchanges with water in the Rio Moquino and Rio Paguate through the unconsolidated alluvium deposits along the perennial river channels. The stream deposits act as a mixing zone between the surface water in the rivers and the ground water discharging from the underlying Jackpile Sandstone Aquifer (Reference 9, Reference 24).

The PPE is the estimated point at which the overland flow segment leading from the source area reaches a perennial surface water body. The PPEs for the Jackpile-Paguate Site are located just north of where the Rio Paguate and Rio Moquino bisect the mine and are in direct contact with Site sources. The PPE locations and surface water pathway are shown in Figure 4-2. From the PPEs, the Rio Moquino and Rio Paguate flow approximately 600 feet before converging and becoming the Rio Paguate. The Rio Paguate flows south approximately 5.5 miles before entering the Paguate Reservoir. The Paguate Reservoir is approximately 0.5 miles long, and then the Rio Paguate continues an additional 0.6 miles before converging with and becoming the Rio San Jose. The Rio San Jose continues for the remaining 15-mile Target Distance Limit (TDL) (Reference 2, Reference 24).

6.2 SURFACE WATER TARGETS

The potential targets of the surface water pathway includes the population relying on surface water within the TDL as a source of drinking water, as well as downstream fisheries, sensitive environments, and surface water resources. No surface water intakes are present within the 15-mile

TDL. Drinking water in the county is provided by ground water. The Village of Paguate obtains water from two wells located in the alluvium of the Rio Paguate upstream from the Site (Reference 3). According to the New Mexico Ecological Services Field Office, there are 8 species of birds, 2 species of fish, 2 species of mammals, 1 anthropod-invertebrate, and 5 plants on the listed and sensitive species list for Cibola County (Reference 15). No commercial fisheries have been identified within the TDL. However, recreational fishing does occur and has been documented within the Rio Paguate, the Paguate Reservoir, and the Rio San Jose. According to the Pueblo of Laguna Environmental and Natural Resources Department, Rio Paguate, Paguate Reservoir, and Rio San Jose are fished for catfish, bluegill, and crappie that are caught for human consumption within the 15-mile TDL (Reference 16).

Surface waters from the Rio Paguate and Rio Moquino within the TDL are used for irrigation upstream from the villages of Paguate and Seboyeta, respectively. Surface water is also consumed by livestock from the Paguate Reservoir and along the Rio Paguate between the reservoir and the Site at points of access (Reference 3). The Rio Paguate's designated uses include cold water aquatic life, domestic water supply, fish culture, irrigation, livestock watering, wildlife habitat, and primary contact (Reference 17).

6.3 SURFACE WATER PATHWAY CONCLUSION

Based on the 2011 ESI surface water and sediment sampling results, a release of isotopic uranium, isotopic thorium, gross alpha and beta, Radium 226 and 228, antimony, arsenic, beryllium, cadmium, cobalt, lead, nickel, and silver to the surface water pathway was documented. The surface water or sediment samples were at concentrations significantly above background levels. ESI sampling locations for the surface water pathway are presented in Table 3-7 and Figure 3-2. Results of the ESI sampling event relating to a release to surface water are presented in Tables 3-8 and 3-9.

The TDL for the Site is 15 miles downstream from the Site in the Rio San Jose. Surface water is in direct contact with the sources within the Jackpile-Paguate Uranium Mine. Further, surface water is used for fishing and livestock and wildlife consumption within the TDL (Reference 24).

7. SOIL EXPOSURE AND AIR MIGRATION PATHWAYS

Other potential routes of hazardous substance migration from the Site include soil exposure or air migration. Factors such as accessibility, likelihood of exposure, and potential targets are evaluated.

No people are known to reside within the site boundaries of the Jackpile-Paguate Uranium Mine site. The nearby population within 4 miles of the site is approximately 500 persons. The site is fenced and gate access is limited. START-3 noted a few locations where the gates and/or fences were in disrepair; however, the Pueblo of Laguna conducts regular inspections to repair the fence. START-3 also noted evidence of cattle grazing at the site (Reference 24).

8. SUMMARY

START-3 conducted an ESI sampling event in April 2011, which included sampling and chemical/radiological analyses of ground water, surface water, and sediments. The specific sampling objectives for the ESI were to collect data that could be used to document a release of hazardous substances to the surface water or ground water pathway. Based on the results of the ESI sampling event, a release to the surface water pathway attributable to the Site was documented.

Based on the 2011 ESI, the following are associated with the Site:

- Estimated quantities of wastes in Site sources include 1,105 acres of material in the open pits, 1,266 acres of material in the waste dumps, and 10.3 million cubic yards of material in the protore stockpiles.
- Approximately 23 million tons of uranium resources remain at the mine site as stockpiled protore and unmined deposits.
- Uranium was detected in Site sources at concentrations significantly above background, up to 364 milligrams per kilogram (mg/kg), during the March 2010 SI sampling. Sources also contained chromium, magnesium, manganese, nickel, selenium, and vanadium at concentrations significantly above background.
- Isotopic uranium, isotopic thorium, gross alpha and beta radiation, Radium 226 and 228, beryllium, cadmium, chromium, cobalt, lead, nickel, selenium, silver, and zinc were all present in ground water at concentrations significantly above background. Observed releases in ground water have been documented in 8 wells.
- Based on the START-3 2011 ESI surface water and sediment sampling results, a release of isotopic uranium, isotopic thorium, gross alpha and beta, Radium 226 and 228, antimony, arsenic, beryllium, cadmium, cobalt, lead, nickel, and silver to the surface water pathway is documented. Releases in surface water have been documented in 7 locations within the surface water pathway TDL. In sediment, radionuclides were documented as being significantly above background at 7 locations, antimony at 2 locations, and cadmium at 1 location. In surface water, radionuclides and metals were documented as being significantly above background at 5 locations.
- The TDL for the Site is 15 miles downstream from the Site in the Rio San Jose. Surface water is in direct contact with the sources within the Jackpile-Paguate Uranium Mine. Surface water is used for fishing and livestock and wildlife consumption within the TDL.

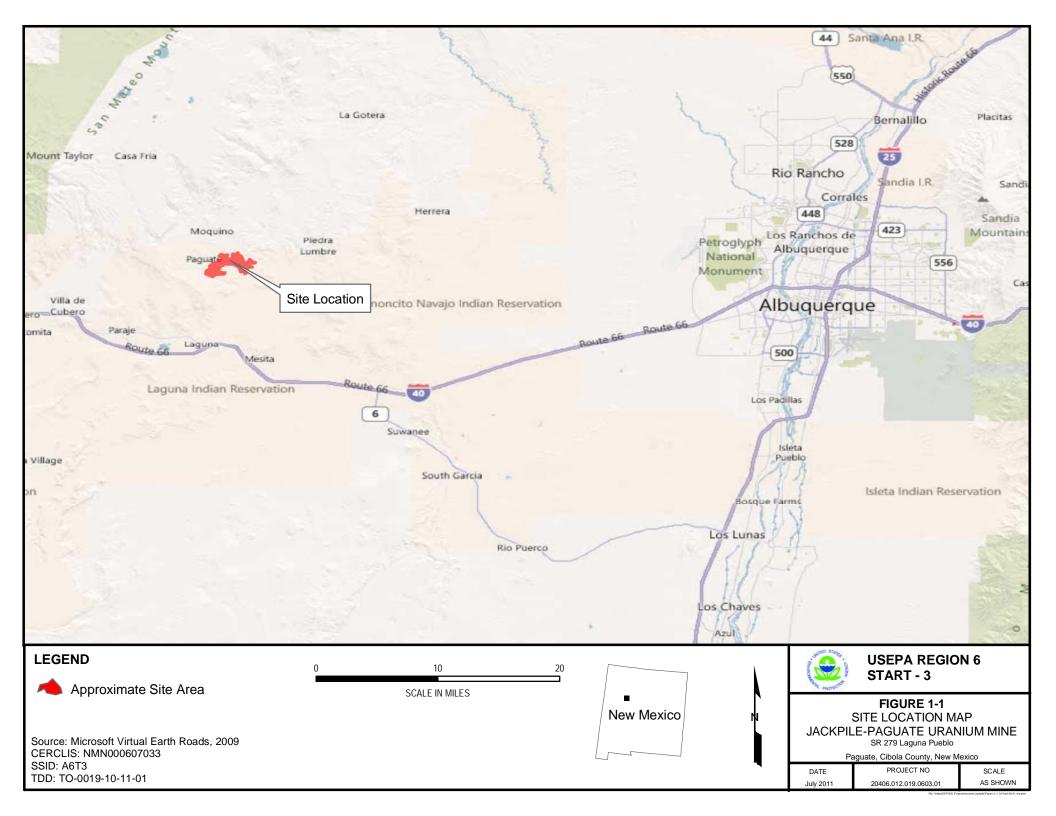
9. REFERENCES

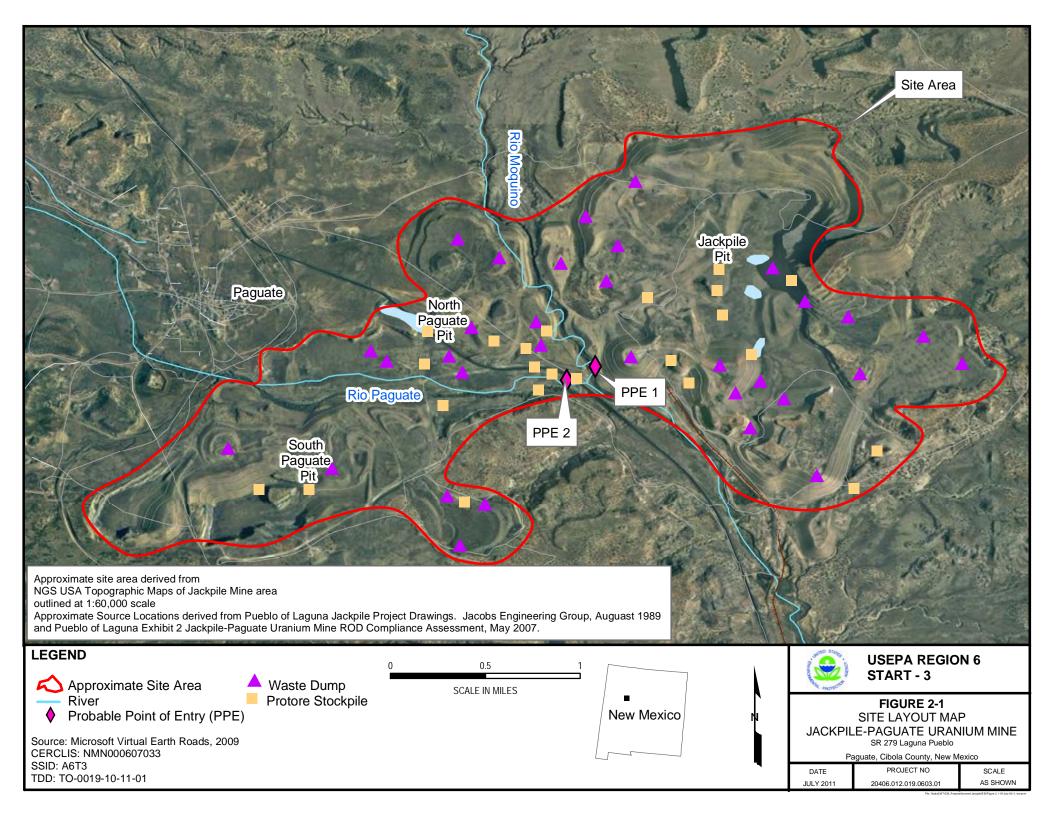
The following reference documents are found as separate files in Appendix G.

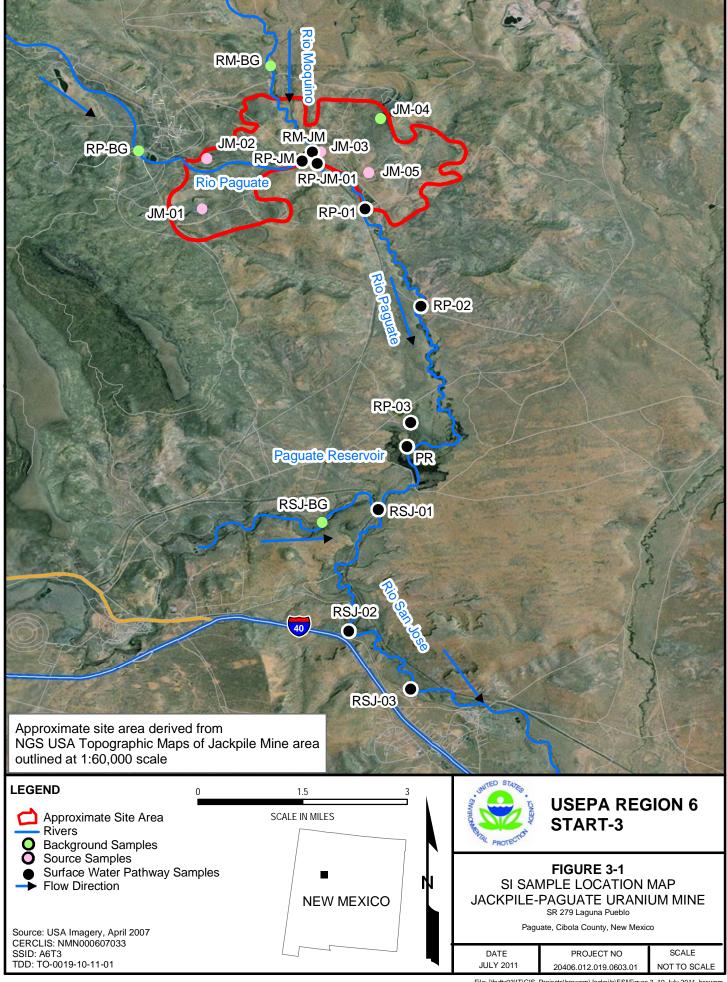
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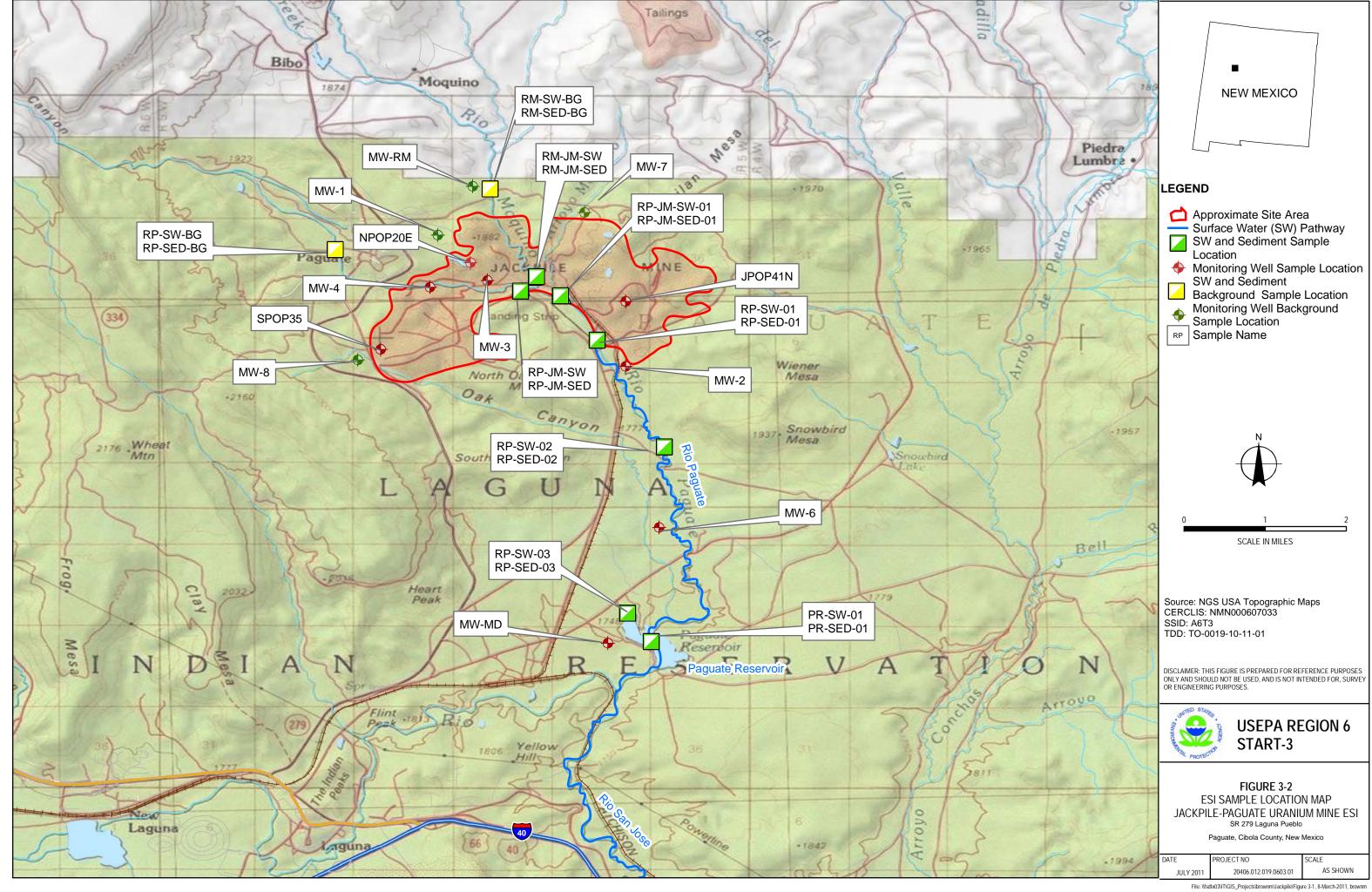
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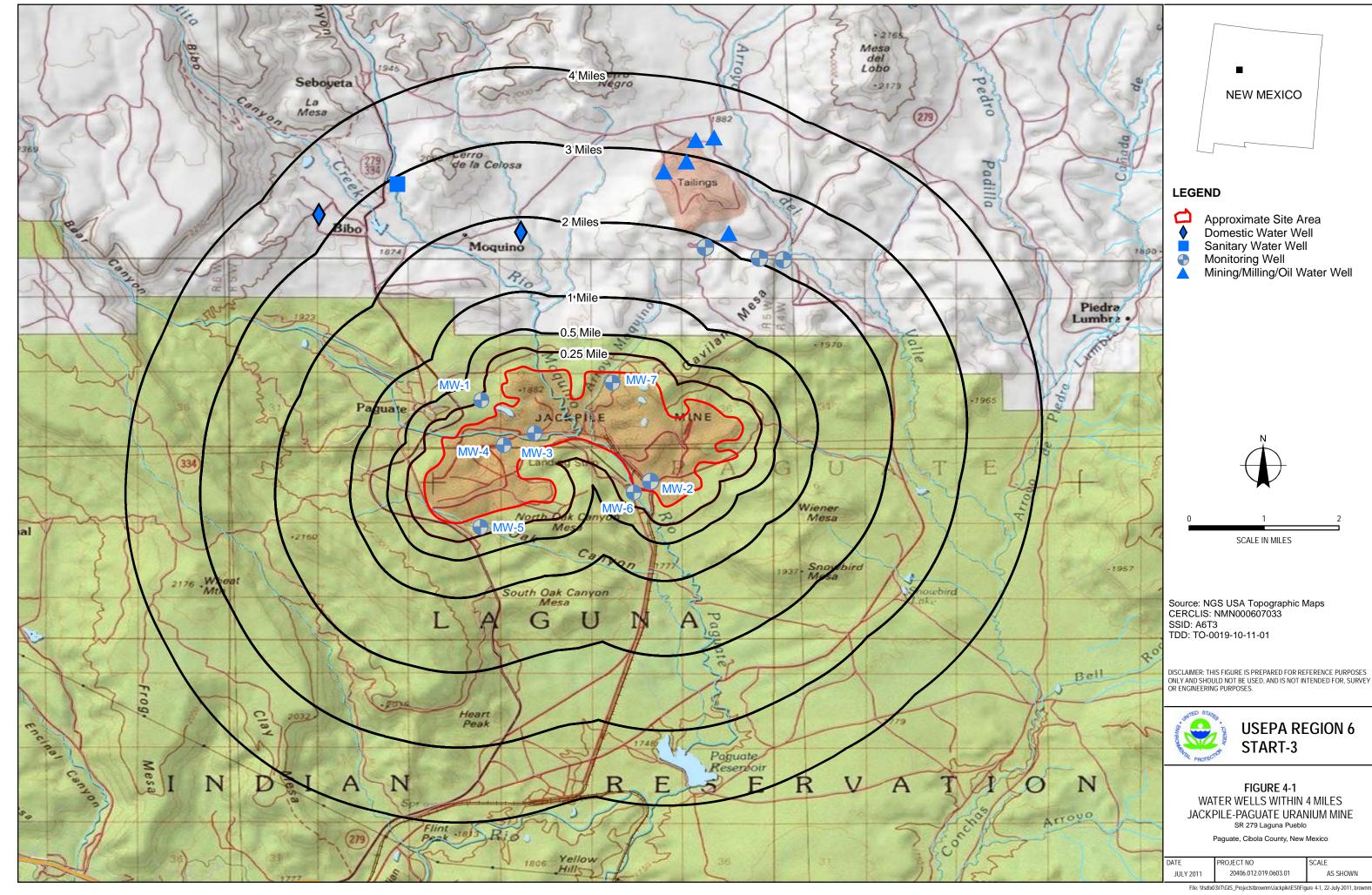
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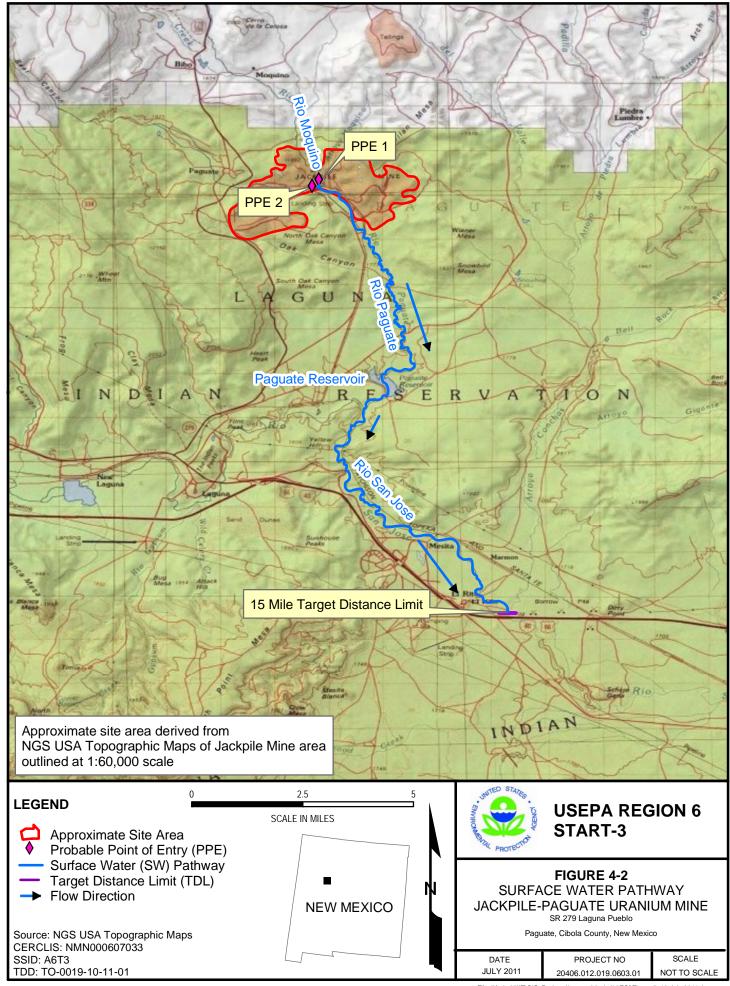


	Table 3-5: Summary of SI Surface Water Pathway Analytical Data - Sediment																
					RP-SED-												
		Residential	RM-SED-BG	RP-SED-BG	BGD	RSJ-SED-BG											
Analyte	Units	Soil RSL	Background	Background	Background	Background	RM-JM-SED	RP-JM-SED	RP-JM-SED	RP-SED-01	RP-SED-01D	RP-SED-02	RP-SED-03	PR-SED-01	RSJ-SED-01	RSJ-SED-02	RSJ-SED-03
U-232	mg/kg	-	2.48E-07	2.51E-07	2.71E-07	2.38E-07	2.6E-07	1.49E-07	2.52E-07	2.28E-07	2.07E-07	2.44E-07	2.71E-07	2.03E-07	2.3E-07	2.89E-07	1.49E-07
U-234	mg/kg	-	0.000162	0.000159	0.000156	0.000069	0.000136	0.000169	0.000238	0.00014	0.00019	0.000227	0.000386	0.00013	0.000039	0.00014	0.00005
U-235	mg/kg	-	0.034	0.028	0.028	0.014	0.019	0.032	0.022	0.021	0.016U	0.033	0.045	0.026	0.0044	0.026	< 0.015
U-238	mg/kg	-	3.53	2.52	2.84	1.42	2.51	3.3	3.9	2.7	3.67	4.67	7.5	2.28	0.74	2.46	0.85
Uranium, Total	mg/kg	230	3.56	2.55	2.86	1.44	2.52	3.33	3.92	2.72	3.69	4.71	7.6	2.31	0.74	2.49	0.86
Aluminum	mg/kg	77000	8400	6800	7300	5400	8100	5900	5000	4700	4300	6400	11000	2400	2100	9800	2800
Arsenic	mg/kg	0.39	6.9	6.8	5.4	2.3	6.5	5.4	4.8	4.3	4.3	5.7	7.9	1.3	1.3	5.2	1.5
Barium	mg/kg	15000	100	88	110	100	75	110	56	66	76	110	130	24	73	150	77
Calcium	mg/kg	1	20000	16000	18000	14000	18000	15000	38000	15000	17000	22000	27000	19000	6900	25000	6200
Chromium	mg/kg	120000 (1)	8.8	6.9	7.7	4.2	8.3	6	7.7	6	4.9	6.4	9.8	1.3	1.5	7.7	2.6
Cobalt	mg/kg	23	6.6	6.5	5.9	2.9	6.4	5.5	5.2	3.8	3.9	5.3	7.3	2.2	1.2	6.1	1.6
Copper	mg/kg	3100	12	7.5	8.7	4.8	12	9.6	7.5	5.9	5.4	9.3	14	2.2	1.5	11	2.4
Iron	mg/kg	55000	23000	24000	20000	8500	21000	21000	17000	13000	14000	17000	24000	3700	3700	18000	5100
Lead	mg/kg	400	12	9.3	9.9		12	9.5	7.2	8.3		11	15	4	2.4	11	2.9
Magnesium	mg/kg	-	5400	4100	4600	2700	5300	4200	4600	3700	3300	5100	6700	2400	1100	5000	1300
Manganese	mg/kg	1800	200	220	270	160	220	240	180	180	200	210	260	500	83	360	75
Nickel	mg/kg	1500	13	11	11	5.2	13	10	11	7	7.3	9.8	14	2.9	2.5	10	3.2
Potassium	mg/kg	1	1800	1300	1600	1200	1700	1200	950	1000	930		2500	450	440	2200	580
Sodium	mg/kg	-	310	280	150	280	370	310	300	190	180	270	470	<u>1500</u>	130	500	160
Vanadium	mg/kg	390	21	23			18	17	18	14	13	16	23	6.1	5	19	6.9
Zinc	mg/kg	23000	46	40	39	18	44	40	29	25	24	36	50	7	7.4	39	10

RSL: June 2011 EPA Regional Screening Table

mg/kg: milligrams per kilogram

BOLD and underlined values are significantly above the background concentration

<: analyte not present at concentration above the associated reporting limit

^{(1) -} The RSL identified for chromium is based on trivalent chromium

	Table 3-6: Summary of SI Surface Water Pathway Analytical Data - Water																
Analyte	Units		RM-SW-BG Background		RP-SW- BGD Background	RSJ-SW-BG Background*	RM-JM-SW	RP-JM-SW	RP-JM-SW-01	RP-SW-01	RP-SW-01D	RP-SW-02	RP-SW-03	PR-SW-01	RSJ-SW-01	RSJ-SW-02	RSJ-SW-03
U-232	μg/l	-	3.82E-07	3.58E-07	3.60E-07	3.68E-07	3.70E-07	3.36E-07	3.61E-07	3.42E-07	3.50E-07	3.21E-07	3.46E-07	3.16E-07	3.49E-07	3.65E-07	3.56E-07
U-234	μg/l	-	0.000341	0.000231	0.000256	0.000474	0.00117	0.00148	0.00126	0.00356	0.00335	0.00343	0.00347	0.0139	0.000466	0.0005	0.000461
U-235	μg/l	-	0.03	0.016	0.028	0.048	0.132	0.16	0.153	0.468	0.39	0.405	0.419	1.75	< 0.022	0.058	0.033
U-238	μg/l	-	3.81	1.78	1.97	3.7	16.7	22.7	<u>18.4</u>	<u>59.8</u>	56.7	56.5	<u>56.6</u>	224	4.28	4.43	4.15
Uranium, Total	μg/l	30	3.84	1.8	2	3.75	16.8	22.9	<u>18.6</u>	60.3	<u>57.1</u>	56.9	<u>57.1</u>	226	4.3	4.49	4.18
Aluminum	μg/l	-	< 200	< 200	200U	770	< 200	400	< 200	<200	200U	370	< 200	< 200	610	720	430
Calcium	μg/l	-	100000	54000	54000	98000	130000	130000	130000	140000	140000	140000	150000	440000	98000	96000	100000
Iron	μg/l	-	180	270	260	530	200	740	330	400	410	610	<100	<100	460	520	320
Magnesium	μg/l	-	56000	19000	19000	53000	90000	52000	74000	93000	91000	90000	91000	230000	53000	52000	55000
Manganese	μg/l	-	27	45	44	56	72	140	95	<u>190</u>	<u>190</u>	87	<10	250	45	42	23
Potassium	μg/l	-	5900	3700	3700	7800	7600	6100	7000	7400	7400	7500	13000	19000	7800	7900	8000
Sodium	μg/l	-	94000	21000	21000	170000	140000	46000	98000	120000	120000	120000	140000	420000	170000	170000	180000

MCL: Maximum Contaminant Level

Bold and underlined values are significantly above background

Highlighted values exceed MCLs

<: analyte not present at concentration above the associated reporting limit

μg/l: micrograms analyte per liter water

^{*} These background concentrations only used for samples collected below convergence with Rio San Jose

	Table 3-8: ESI Sampling Results for Radionuclides												
Location	Location Description		Th ²³⁰	Th ²³²	U^{234}	U^{235}	U^{238}	Utot	Thtot	Gross a	Gross β	Ra ²²⁶	Ra ²²⁸
Groundwater		pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l
MW-1	Background	0.043963821	0.122151363	0.007171001	3.743294543	0.160700599	0.825715965	4.730	0.173	13.4	< 3.58	< 0.74	1.26
MW-7	Background	0.035638155	0.485386768	0.038753264	3.80344807	0.061541945	0.44316914	4.308	0.560	1047	<3.58	2.71	1.11
MW-8	Background	0.004018533	0.937884647	0.011798427	5.209553086	0.058512416	1.954712573	7.223	0.954	40.5	224	< 0.74	< 0.92
MW-RM	Background	0.012318119	1.034526789	0.104450935	11.64534003	0.732724784	30.02396421	42.402	1.151	<2.28	11.22	< 0.74	< 0.92
NPOP20E	Source	0.037393028	1.853608356	0.109895077	<u>36513.126</u>	40367.1181	64338.356	141218.600	2.001	<u>3624</u>	<u>26301.04</u>	33.23	13.93
SPOP35	Source	0.0754155	0.592061916	0.09971059	7837.898207	6449.591318	9952.40578	24239.895	0.767	1326	<u>2314.11</u>	<u>17.29</u>	<u>17.3</u>
JPOP41S	Source	0.247162371	1.491832528	0.086560951	63261.3878	72121.57278	114131.786	<u>249514.747</u>	1.826	<u>15961</u>	147252.19	208.45	38.3
MW-2	On-Site	0.012825277	0.004200614	-0.071267598	63.555318	2.232365086	54.37424512	120.162	-0.054	65.5	99.3	< 0.74	< 0.92
MW-2-2	On-Site	0.008448284	0.041505515	-0.004142249	62.70805081	2.489819913	54.41057754	119.608	0.046	70	120.3	< 0.74	< 0.92
MW-3	On-Site	0.007967807	0.645251517	0.031222341	<u>17.71281407</u>	0.649610379	15.43632023	33.799	0.684	43	5.3	< 0.74	< 0.92
MW-4	On-Site	< 0.013	0.128568173	0.008019437	118.9504898	4.57933476	108.2253186	231.755	0.137	142	91.3	< 0.74	< 0.92
MW-6	Downgradient	0.026390062	0.673519252	0.040625778	73.20502395	2.628962984	61.7977617	137.632	0.741	61.8	66.7	< 0.74	2.51
MW-MD	Downgradient	0.009227451	0.858537855	0.008698685	31.28938758	1.000809104	22.77285666	<u>55.063</u>	0.876	27	17.8	1.34	< 0.92
MW-MD-02	Downgradient	-0.004057242	0.692972714	0.023847811	29.05916125	1.065731024	21.35722039	<u>51.482</u>	0.713	149	20.7	< 0.74	< 0.92
	Surface Water	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l	pCi/l
RM-SW-BG	R. Moquino Bkgd	0.071110945	0.72692127	0.12597932	4.119181833	0.099619372	2.230079254	6.449	0.924	<2.28	5.6	< 0.74	1.12
RP-SW-BG	R. Paguate Bkgd	0.161190066	1.202221671	0.072716165	6.241783844	0.215676269	3.79272602	10.250	1.436	<2.28	<3.58	< 0.74	< 0.92
RM-JM-SW	R. Moquino above confluence	0.036489552	0.544542865	0.138838501	<u>37.99668754</u>	<u>1.540503892</u>	<u>33.43142295</u>	<u>72.969</u>	0.720	<u>28.2</u>	<u>45.6</u>	< 0.74	< 0.92
RM-JM-SW-02	R. Moquino above confluence	0.027277183	0.871007328	0.110564403	<u>28.33431683</u>	0.960622987	<u>23.59076426</u>	<u>52.886</u>	1.009	<u>23.6</u>	<u>39.4</u>	< 0.74	2.02
RP-JM-SW	R. Paguate above confluence	0.226013156	0.702348426	0.141857603	28.47649254	1.070865743	<u>24.24662061</u>	53.794	1.070	<u>35.1</u>	<u>23.7</u>	< 0.74	< 0.92
RP-JM-SW-01	R. Paguate below confluence	0.026113869	0.755229107	0.076649323	32.3975508	1.235555446	29.25685624	<u>62.890</u>	0.858	<u>30.8</u>	<u>25.9</u>	< 0.74	< 0.92
RP-SW-01	R. Paguate site south site boundary	0.104096263	1.072519788	0.109886334	<u>85.09851669</u>	3.235728052	<u>78.78713513</u>	<u>167.121</u>	1.287	<u>75</u>	<u>28.1</u>	< 0.74	< 0.92
PR-SW-01	Paguate Reservoir	0.033888613	1.572556901	0.08841753	<u>17.44528511</u>	0.564714659	14.62398112	32.634	1.695	<u>55</u>	<u>151.8</u>	< 0.74	< 0.92
	Sediment	pCi/g	pCi/g	pCi/g	pCi/g	pCi/g	pCi/g	pCi/g	pCi/g	pCi/g	pCi/g	pCi/g	pCi/g
RM-SED-BG	R. Moquino Bkgd	0.821	1.045	0.821	0.369	0.020	0.303	0.692	2.687	4.8	4.1	1.17	<2
RP-SED-BG	R. Paguate Bkgd	0.465	0.454	0.485	0.243	0.002	0.202	0.446	1.403	1.9	1.34	1.71	<2
RM-JM-SED	R. Moquino above confluence	<u>1.266</u>	<u>6.829</u>	<u>1.268</u>	<u>5.781</u>	<u>0.262</u>	<u>5.888</u>	<u>11.931</u>	<u>9.363</u>	5.45	<u>9.75</u>	<1	<2
	R. Moquino above confluence	0.548	0.973	0.452	0.820	0.054	0.846	<u>1.720</u>	1.973	2.43	<3	1	2.64
RP-JM-SED	R. Paguate site south site boundary	1.006	0.859	0.908	<u>0.786</u>	0.029	<u>0.691</u>	<u>1.506</u>	2.773	4.6	2.4	1.77	<2
	R. Paguate above confluence	0.371	0.679	0.419	0.680	<u>0.049</u>	0.688	<u>1.417</u>	1.469	2.8	1.99	<1	<u>2.14</u>
RP-SED-01	R. Paguate below confluence	0.512	0.584	0.476	<u>0.711</u>	0.015	0.631	<u>1.356</u>	1.573	1.5	1.13	1.75	<2
RP-SED-02	R. Paguate downgradient	0.795	<u>1.610</u>	0.733	0.903	0.031	<u>0.947</u>	<u>1.881</u>	3.138	5.44	4.8	1.93	<2
RP-SED-03	R. Paguate downgradient	1.130	3.982	0.995	<u>1.350</u>	<u>0.101</u>	<u>1.624</u>	<u>3.076</u>	<u>6.107</u>	4.5	3.4	<u>2.73</u>	<2
PR-SED-01	Paguate Reservoir	0.462	4.433	0.543	2.124	<u>0.089</u>	<u>2.247</u>	4.460	<u>5.437</u>	4.83	<u>9.6</u>	<u>3.17</u>	3.4

pCi/l: picoCuries per liter pCi/g: picoCuries per gram

<: analyte not detected above the associated reporting limit bold and underlined values are significantly above background

	Table 3-9: ESI Sampling Results for Metals																
	Designation	Mecury	Antimony	Arsenic	Barium	Beryllium	Cadmium	Chromium	Cobalt	Lead	Manganese	Nickel	Selenium	Silver	Thallium	Vanadium	Zinc
G	Froundwater	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l
MW-1	Background	< 0.00006	0.0075 UB	< 0.00206	0.018 JQ	< 0.00072	< 0.00082	< 0.00109	0.0022 UB	<0.0021 JL	< 0.00043	0.0025 UB	0.0309 UB	0.0014 UBJL	0.0212 UB	< 0.00174	< 0.00053
MW-7	Background	< 0.00006	0.003 UB	0.0028 UB	0.0274 JQ	< 0.00072	< 0.00082	< 0.00109	0.0006 UB	0.0029	< 0.00043	< 0.00116	0.0296 UB	< 0.00039	0.0119 UB	< 0.00174	0.002 UB
MW-8	Background	< 0.00006	0.0096 UB	0.0033 UB	0.0389 JQ	< 0.00072	< 0.00082	0.0012 UB	0.001 UB	< 0.0021	< 0.00043	0.0019 UB	0.0317 UB	0.0011 UBJL	< 0.00529	< 0.00174	< 0.00053
MW-RM	Background	< 0.00006	0.0114 UB	< 0.00206	0.0134 JQ	< 0.00072	< 0.00082	< 0.00109	< 0.00051	< 0.0021	< 0.00043	< 0.00116	0.0344 UBJL	< 0.00039	0.0082 JQ	< 0.00174	< 0.00053
NPOP20E	Source	< 0.00006	< 0.00237	< 0.00206	0.0112 JQ	0.0067	< 0.00082	0.0604		0.0519 UBJL	4.633	0.0374 UB	0.0309 UB	0.0522 JL	< 0.00529	< 0.00174	0.0107 UB
SPOP35	Source	< 0.00006	< 0.00237	< 0.00206	0.007 UB	< 0.00072	< 0.00082	0.0057 JQ	0.0016 UB	0.008 UBJL	0.186	0.0062 UB	0.0337 UB	0.0047 UBJL	< 0.00529	< 0.00174	< 0.00053
JPOP41S	Source	< 0.00006	< 0.00237	< 0.00206	0.008 UB	0.0395	0.0045 UB	<u>0.365</u>	0.0788	0.341 JL	<u>2.84</u>	0.0086 UB	0.104 UB	0.321 JL	< 0.00529	< 0.00174	0.0155 UB
MW-2	On-Site	< 0.00006	0.0061 UB	0.0033 UB	0.0207 JQ	< 0.00072	< 0.00082	< 0.00109	0.0006 UB	0.003 UBJL	0.275	0.0048 UB	0.0317 UB	0.002 UBJL	0.0106 UB	< 0.00174	0.0017 UB
MW-2-2	On-Site	< 0.00006	0.0057 UB	0.00206 UB	0.0221 JQ	< 0.00072	< 0.00082	< 0.00109	0.0006 UB	<0.0021 JL	0.294	0.0052 UB	0.0327 UB	<0.00039 R	0.0077 UB	< 0.00174	< 0.00053
MW-3	On-Site	< 0.00006	0.0024 UB	< 0.00206	0.0281 JQ	< 0.00072	< 0.00082	< 0.00109	< 0.00051	< 0.0021	< 0.00043	< 0.00116	0.0337 UB	<0.00039 R	< 0.00529	< 0.00174	0.0066 UB
MW-4	On-Site	< 0.00006	0.0207 UB	0.0041 UB	0.0357 JQ	0.0008 UB	< 0.00082	< 0.00109	0.0011 UB	0.0027 UBJL	0.0324	0.0022 UB	0.0395 UB	0.0025 UBJL	0.0075 UB	< 0.00174	0.0103 UB
MW-6	Downgradient	< 0.00006	0.0056 UB	< 0.00206	0.0148 JQ	< 0.00072	< 0.00082	< 0.00109	< 0.00051	<0.0021 JL	0.0193 UB	0.002 UB	0.0288 UB	0.001 UBJL	0.0081 UB	< 0.00174	0.0037 UB
112 11 1122	Downgradient	< 0.00006	0.0041 UB	0.0064 UB	0.0116 JQ	< 0.00072	< 0.00082	< 0.00109	0.0013 UB	<0.0021 JL	0.654	0.0048 UB	0.0271 UB	<0.00039 R	< 0.00529	< 0.00174	< 0.00053
MW-MD-02	Downgradient	< 0.00006	0.0074 UB	0.0102 UB	0.0132 JQ	< 0.00072	< 0.00082	0.0014 UB	0.0017 UB	<0.0021 JL	0.693	0.0057 UB	0.0395 UB	<0.00039 R	0.0099 UB	< 0.00174	< 0.00053
Su	Surface Water		mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l		mg/l	mg/l	mg/l	mg/l	mg/l	mg/l
RM-SW-BG	R. Moquino Bkgd	< 0.00006	0.0043 UB	< 0.00206	0.0261 JQ	< 0.00072	< 0.00082	< 0.00109	< 0.00051	<0.0021 JL	< 0.00043	< 0.00116	0.035 UB	<0.00039 R	0.0151 UB	< 0.00174	0.0023 UB
RP-SW-BG	R. Paguate Bkgd	< 0.00006	< 0.00237	< 0.00206	0.0742	< 0.00072	< 0.00082	< 0.00109	< 0.00051	<0.0021 JL	0.641	< 0.00116	0.0273 UB	<0.00039 R	0.0111 UB	< 0.00174	0.0046 UB
RM-JM-SW	R. Moquino upper	< 0.00006	0.0041 UB	< 0.00206	0.0228 JQ	< 0.00072	< 0.00082	< 0.00109	< 0.00051	0.0025 UBJL	0.0131 UB	< 0.00116	0.0324 UB	<0.00039 R	0.0175 UB	< 0.00174	< 0.00053
RM-JM-SW-02	R. Moquino upper	< 0.00006	0.0062 JQ	< 0.00206	0.0232 JQ	< 0.00072	< 0.00082	< 0.00109	< 0.00051	<0.0021 JL	0.0152 JQ	0.0023 JQ	0.0278 UB	<0.00039 R	0.0154 UB	< 0.00174	< 0.00053
RP-JM-SW	R. Paguate upper	< 0.00006	0.0048 UB	< 0.00206	0.0472	< 0.00072	< 0.00082	< 0.00109	< 0.00051	<0.0021 JL	0.0374	0.0019 UB	0.0276 UB	0.0012 UBJL	< 0.00529	< 0.00174	0.0034 UB
	R. Paguate lower	< 0.00006	< 0.00237	< 0.00206	0.037 JQ	< 0.00072	< 0.00082	< 0.00109	< 0.00051	<0.0021 JL	0.0348	< 0.00116	0.0341 UB	<0.00039 R	0.0071 UB	< 0.00174	0.0025 UB
RP-SW-01	R. Paguate S. boundary	< 0.00006	0.0047 UB	< 0.00206	0.0255 JQ	< 0.00072	< 0.00082	< 0.00109	0.0007 UB	<0.0021 JL	0.97	0.0025 UB	0.0299 UB	<0.00039 R	0.0082 UB	< 0.00174	< 0.00053
PR-SW-01	Paguate Reservoir	< 0.00006	0.0033 UB	0.0075 UB	0.14	< 0.00072	< 0.00082	< 0.00109	0.0009 UB	<0.0021 JL	0.0424	0.0032 UB	0.0302 UB	<0.00039 R	0.013 UB	< 0.00174	< 0.00053
	Sediment	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg		mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg
RM-SED-BG	R. Moquino Bkgd	< 0.0563	<0.113 JL	6.68	154 JH	0.916 UB	< 0.0282	8.4	7.78	4.58 JQ	226	14.6 JL	29.1	< 0.141	0.662 UB	29.8	57
RP-SED-BG	R. Paguate Bkgd	< 0.0532	<0.106 JL	4.53	97.7 JH	0.586 UB	< 0.0266	9.12	9.34	1.72 UB	358	9.17 JL	28.3	< 0.133	< 0.213	29.9	38.7
RM-JM-SED	R. Moquino upper	< 0.0474	23.3 JK	5.72	57 JH	0.558 UB	< 0.0237	9.1		2.35 UB	173	9.18 JL	23.3 JK	< 0.119	0.717 UB	22.4	36
RP-JM-SED	R. Paguate S. boundary	< 0.0525	<0.105 JL	5.73	72.9 JH	0.729 UB	< 0.0263	11.7	5.63	3.15 UB	166	11.1 JL	23.6	< 0.131	< 0.21	24.3	52.3
RP-JM-SED-02	R. Paguate S. boundary	< 0.0436	0.529 UBJK	4.65	68 JH	0.393 UB	0.24 JQ	7.88	5.22	3.06 UB	160	9.36 JL	<0.109 JK	< 0.109	< 0.174	19.8	34.2
RP-JM-SED-01	R. Paguate upper	< 0.0468	<0.0937 JL	6.57	116 JH	0.615 UB	< 0.0234	9.33	6.29	1.9 UB	490	10.6 JL	30	< 0.117	< 0.187	23.8	34.1
RP-SED-01	R. Paguate lower	< 0.0511	<0.102 JL	4.56	76.2 JH	0.574 UB	< 0.0255	8.09	4	3.32 UB	150	7.54 JL	18.4	< 0.128	0.747 UB	18.6	32.7
RP-SED-02	R. Paguate downgradient	< 0.0515	<0.103 JL	4.52	157 JH	0.728 UB	< 0.0258	9.58	4.88	2.59 UB	258	8.95 JL	19.9 UB	< 0.129	< 0.206	22.2	38.8
RP-SED-03	R. Paguate downgradient	< 0.0419	<0.0838 JL	6.05	132 JH	1.14 UB	< 0.0209	16.1	7.17	3.22 UB	231	14.7 JL	28.7	< 0.105	< 0.168	34.6	60.1
PR-SED-01	Paguate Reservoir	< 0.0632	1.88 UBJL	7.82	187 JH	1.19 UB	<u>0.901</u>	17.7	7.3	9.69	558	14.7 JL	< 0.158	<0.158 R	< 0.253	44.8	65

mg/l: milligrams per liter

mg/kg: milligrams per kilogram

<: analyte not detected above the associated reporting limit

bold and underlined values are significantly above background

UB - undetected due to blank contamination

J - the value is an estimated quantity

L - Low bias

H - High bias

K - Unknown bias

Q - the reported concentration is less than the sample quantitation limit

R - quality control indicates the data are unusable for all purposes

APPENDIX A DIGITAL PHOTOGRAPHS

Jackpile-Paguate Uranium Mine ESI Photo Summary 19 - 21 April 2011



19 April 2011, 11:52 – N - MW-7 location. Pumping purged water.



19 April 2011, 15:39 – N - MW-1 location.



19 April 2011, 1339 - E - MW-8 location. Putting tubing/pump out of well and preparing for bailing.



19 April 2011, 16:54 – S - MW-RM location. Bailing.



19 April 2011, 17:54 – SE - MW-MD location. Southern most monitoring well.



20 April 2011, 12:24 – SE - MW-2 location.



20 April 2011, 11:45 – NW - MW-6 location.



20 April 2011, 13:12 – NE - MW-4 location.



20 April 2011, 14:03- NW - MW-3 location.



20 April 2011, 15:47 – NNE - NPOP20E location.



20 April 2011, 14:48 – N - SPOP35 location in South Pit.



20 April 2011, 16:28 – NW - JPOP41S location.



21 April 2011, 08:14 – S - Yellow groundwater in drinking bottle from JPOP41S collected 20 April 2011.



21April 2011, 09:39 – E - JPOP41S groundwater sample.



21 April 2011, 09:39 – E - JPOP41S groundwater sample.



21April 2011, 09:39 – NW - Morris Formation containing the native uranium below the black strata graphic layer in the crushing area of Jackpile.



21 April 2011, 10:00 –NW - Close-up of the native uranium in crushing area of Jackpile.



19 April 2011, 10:31 – NE - From bridge at brush/location RP-BG.



21 April 2011, 10:00 – NW - Close-up of the native uranium in crushing area of Jackpile.



19 April 2011, 10:31 – SE - From bridge downstream in flow direction. RP-BG location.



19 April 2011, 10:42 – Showing water clarity at location RP-BG.



19 April 2011, 11:13 – Sample location RM-BG looking upstream.



19 April 2011, 11:13-RP-SW-BG.



19 April 2011, 14:32– Sample location RM-BG looking downstream.



19 April 2011, 14:32– Sample location RM-BG overview.



19 April 2011, 14:58– RM-SED-BG.



19 April 2011, 14:57– RM-SW-BG.



19 April 2011, 15:43 - Sampling location RP-01 looking downstream.



19 April 2011, 15:49 - Sampling location RP-01 looking upstream.



19 April 2011, 16:27 – Sample RP-SW-01.



19 April 2011, 16:27 – Sample RP-SW-01.



20 April 2011, 09:32 – S - Facing downstream at RP-02.



20 April 2011, 09:34 - N – Facing upstream at RP-02.



20 April 2011, 10:18 – S - Facing downstream at RP-03.



20 April 2011, 09:36 - RP-02-SED.



20 April 2011, 10:18 – N - Facing upstream at RP-03.



20 April 2011, 10:23 – RP-03-SED.



20 April 2011, 10:57 – Wildlife at Reservoir – snake.



20 April 2011, 10:25 –E – Overview of sample location RP-03.



20 April 2011, 11:38 – Wildlife at Reservoir – frog.



20 April 2011, 11:38 - SW- Pooled water within Reservoir bank.



20 April 2011, 11:49 - Wildlife at Reservoir – another snake.



20 April 2011, 11:38 – N - Overview of sample location PR.



20 April 2011, 11:50 – NW - Overview of sample location PR.



20 April 2011, 11:50 - Different variety of snake present at Reservoir.



20 April 2011, 14:04 – Overview of the Mesita Diversion.



20 April 2011, 11:58 – Samples PR-SED and PR-SW.



20 April 2011, 14:04 – Overview of the Mesita Diversion.



20 April 2011, 14:18 – Mesita Diversion (MD) surface water sample.



20 April 2011, 15:50 – NW- Facing PR and PM confluence.



20 April 2011, 15:50 – SE - Facing downstream sat RP-JM-01.



20 April 2011, 15:50 – Overview of RP-JM-01 sample location.



20 April 2011, 16:11 – RP-JM-01-SW



20 April 2011, 16:12 – S - Rio Paguate through conduit.



20 April 2011, 16:12 – RP-JM-01-SED



20 April 2011, 16:12 – Rio Paguate flow out at mouth of conduit..



20 April 2011, 16:12 – S – Channel and sediment of Rio Paguate.



20 April 2011, 16:33 – RP-JM-SW and RP-JM- SED samples.



20 April 2011, 16:16 – SW – START-3 collecting surface water sample.



20 April 2011, 16:57 – E - START-3 measuring Rio Moquino flow.



20 April 2011, 16:58 – E - Facing downstream on Rio Moquino facing confluence.



20 April 2011, 17:34 – RM-JM-SW (normal).



20 April 2011, 16:58 – E - Facing upstream on Rio Moquino.



20 April 2011, 17:37 – RM-JM-SED (normal).



20 April 2011, 17:37 – RM-JM-SW-02 (duplicate).



21 April 2011, 16:15 – Rio Paguate, upstream of Rio Paguate and Rio Moquino confluence – an exposed eroded face > 999uR/hr



20 April 2011, 17:40 – RM-JM-SW-02 (duplicate).



21 April 2011, 16:15 – Rio Paguate, upstream of Rio Paguate and Rio Moquino confluence – an exposed eroded face > 999uR/hr



21 April 2011, 16:27 – Rio Paguate, upstream of Rio Paguate and Rio Moquino confluence - material leaching from bank



21 April 2011, 16:28 – Rio Paguate, upstream of Rio Paguate and Rio Moquino confluence - material leaching from bank



21 April 2011, 16:27 – Rio Paguate, upstream of Rio Paguate and Rio Moquino confluence - material leaching from bank



21 April 2011, 16:29 – Rio Paguate, upstream of Rio Paguate and Rio Moquino confluence - showing efflorescent on bank



21 April 2011, 16:31 – Rio Paguate, upstream of Rio Paguate and Rio Moquino confluence - high emitting (<999 uR/hr) material on bank



21 April 2011, 16:29 – Rio Paguate, upstream of Rio Paguate and Rio Moquino confluence ->999 uR/hr deposit on boulder in channel

APPENDIX B START-3 SITE LOGBOOKS



Jackpile EST Groundwater Sampling Team TO-0019-10-11-01-A

"Rite in the Rain" ALL-WEATHER WRITING PAPER

Name Jackpile ESI

Weston Solutions

Address 14120 N Dallas Pkwy

Dallas TX 75254

Phone 4169. 374.7700

Project Jackpile ESI

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	CONTENTS	* >2
PAGE	REFERENCE	DATE
	Patrick Buster	
	Patrick Buster Tori Gomen & To	5
	E.	

11.19.140 TOD TO-0019-10-11-01-A Personnel: PTL Michelle Brown, FSO James Beauts & Tori Buster, ¿ Dean Frongling (Weston) Physical: Trips slips & Rolla, Turain usi sun Dreem, Chap stick, cactus, Wind Remote Areas-stay with team Biological: Wild animab - bears maintains lions, snakes poisionais non, cattle. Plants - cactus. Weather: Varible cloudiness and windy Hit HITS. Winds W 20-30 mph. Occasional quest over 40 mph. Hosiptal: Acoma Service Unit west on I-40 ~ 24-25 mi. 1000 Groundwater sampling team PBuster & Towney depart Natural Resources Rectamation to meet up with Elvin Garcia, Summit Emiranental and Marvin. 1010 Ground water Sampling Team arrives at meeting location. Summit Environmental and bun Castellana (Weston) on situ. Weston waiting for permits to be signed. [1040 Michelle Billion (Western) and EPA SAM Ladonna Turner arrived with permits.

04.19.11 TOD-TO-0019-10-11-01-A 1047 Groundwater Sampling + departed theter Ground of MW-7 4405 Team includes Weston, Summit Environmental and Marvin . -IIID BWST arrive at MW-7 and begin Set up for purging and sampling. Equipment: GPS ALGML Camera A69935. GPS coordinates ? N 35,15290 W 107, 34133 - 11ft 1120 Summit Environmental puts Redi-flo pump into MN-7/110419 1125 Bigin pumping at MW-7-100 10g/nin Equipment: Ludlum 192 #249765 Water Level Meter #17770, Geopump #21148. MW-7 Parameters. LL Reading 8 micro-R/hc. Time of Temp Conduct DOlo DOM LORP gallins Unital 8,40 21.93 878 21.41 1.79 HOT 1155 8.50 18.87 858 31.6 2.42 lele.9 8.53 18,29 850 21.1 2.53 77.1 1215 8,54 18,10 850 26.5 2.35 57.9 1225 8.55 18.08 847 31,8 3.08 lele-1235 8.45 819 17,73 31.8 2.94 71.0 tiltes correct sample volume later.

TOO TO-BB19-10-11-91-A for departure. 1252 GWST depart to MW-8-110419 1310 GWST arrived at MW-8-110419 location Beging -1315 Begin set up of equipment. LL Reading: 8 micro-R/hr. 1328 Kump controller box gave an evor message. Will not be able to pump MW-8 with equipment. Pumpour heated, Will Bail out 3g of water and take parameters. Prepare For bailing. Pull bump and dean GPS: N 35,12472 W107,3885 +9H 1347 begin bailing MW-8-110419 time of Temp Cord 00% Dongle unihal 8.05 18.97 1429 58.1 5,25 1417 8.02 18,72 1658 60.8 1430 1.97 17.83 1656 57.0 5.33 98.2 1452 8.00 1794 1656 60.1 5.64 1452 Collected 39 of water from \$ for MW. 8-110419. Will Letter and collect correct sample volume later. Decon equipment. 1515 MBrown Blastellana arrived at

64.19.11 TOO TO-0019-10-11-01-A and PBuster explained to mBrown about the parameter & sampling Completed at MW-8-110419. -1525 GNST departed to MW-1-110419 1533 Arrived at MW-1. Begin Set up of equipment. 1540 Bigin Dutaping at mw-1, 10g/min W 107.36750 = 18 Galloro 6PS N 35, 14049' time of Temp 00,ng/L ORP TOWNE Mittel 7.66 18,49 2581 26.1 2.39 90.9 Hofming 1546 7.85 16.82 2478 32.2 3.00 93.5 155/ 7.86 16.76 2428 31.5 2.14 105,2 100 2,14 150 1601 7.79 14.77 2420 17.7 1.68 1606 7.85 16.60 2-154 26.5 2.46 120.0 250 1611 7.78 16.76 2597 34.6 3,72 133.6 300 MW-1-110419. Will filter and collect Correct sample volume later, equipment. Duplicate sample & MS/MSD 1633 GWST departed for MN 1644 Arrived at MW-RM-110919 1646 Begin bailing MW-RM-110419 GPS Nº 35. 15302 V LL: 6.5 micro-R/hr. W107.35631 = 17H

04.19.11

Photolog.

11.91.10

64.19.11

04,19,11

04,9.11 Enviro bailing.
MW-MO location, Southern
most monitoring well. SE 104.19.11

8 TOO TO-18019-115-11-01-A temp Cond DO % Domy/2 ORP gallons 94.2 15,21 9.27 14,42 1919 9.85 115.6 905 14.10 96.7 9.85 102.8 13.89 98.2 10.03 121. 14.02 1916 97.1 10297159 impleted bailing MW-RIA went dry. Will sample 54.20.11. Departed for MW-MD - 110419 1745 Anived at MW-MD-110419 equipmen 1753 Begin pumping a DOTO DOMO L ORP gallons 23.35 91.8 7.71 51.2 6995 7.46 16.40 47.9 4.57 7.40 15.43 7005 21.1 1.96 -124.8 1800 1802 MW-MD - 110419 pumped dry, Will 04.20.11. - reading: 7 micro- R/hc. 1817 GWST departed to command post 1835 GWST arrived of command post. Unload equipment. 1850 Denkob to

TDO TO-0019-10-11-01-A 04.20.11 at MW-6. Decon equipm and prepare for departur 1215 GWST departed for MW-2 222 Arrived at MW D2 location to purge, take parameters GPS: N35,12388 W 107,33591 INF LL Reading: 20 micro-R/hr 1226 begin pumping MW-2.59/min time pH temp Cond ODO OUNGLORP Gal unitial 8.62 20.75 41 199.6 10,98 7,77 -38.2 20 1229 7.34 16.6 3617 63.8 7.38 16.20 492 -905 25 3689 1290 MW-2 went dry, wait for licharge 1250 begin against pumping again 1252 No water, GWST will return later to sample. Decon equipment. 1257 MW-2 has a lot of sediment within water then what has occurred at this will from previous years. Summit Emvorment stated 1302 GWST departs to MW-4 1306 AMIRED at MW- 4. begin set up. with bailers. 685: N 35.18336. W 107.35591 " INGT.

TOD TO-0019-10-11-01-A LL reading: 12 mirc-R/hr. at 59/min time pH temp Cond 100% Domy inital 8,86 21.61 50 131.1 11.56 1402 7.27 14.30 1215 10.54 10 108.7 OLitas 1404 7.14 12.79 1307 78.3 10.17 1406 7.01 12.54 1305 94.0 9.85 56.3 1409 7.14 12.39 1305 95.0 9.99 58.4 1413 6.99 12.29 1306 99.7 10.62 1299 95.0 1415, 6.95 11.96 10.03 Jample MW-3-110420 1415 Collected 500ml bottles at MW-3 supment and prepare 4430 GWST departs on SPOP 35 -1442 GWST arrives at Begin set up. N 35 6PS N 35.12493 N107.38263 = 214 LL reading: 12 micro-R/hr. 1449 Put pump down 590935. 1450 Buyin pumping at SPOP35 at 5g/min. Parameters on

18				
04.20.11 T	DO TU-	-0019-	10-11-01-A	
1710 GWST	1710 GWST depart JPOP415 for			
armand	post.	START.	-3 Phister	
and Bun	es Che	cked	upon	
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Leavene &	comm	and	00st. 5WST	
· sampline	at r	ver 1	fork where	
Rio Paquate	and A	lo M	oguino meet.	
1715 SWST	Thas 1	ust 1	inished and	
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04.21.11 TOD TO-0019-10-11-01-A 0730 START-3 PBuster and Tromes and EPA SAM Ladonna Turner · arrive at spill way, enterance to gockpile open pit mine to meet up with Marvin Saraccino; Laguna tribe member and Marvin Chavery with Summit Environmented to begin sampling on JPOP41S. 0739 Bigin calibration of YSI#24949 (Phister performing calibration) Atmospheric prissure: 604.7mmHg DOJO -> 79.9% @ 17.28°C-Conductivity 1000 pcs/cm > 1000 @ 18.59℃ ORP 228 mm > 227.9 DH 10 7 10.00 @ 17.71°C DH 4 -> 4,00 @ 17,00°C pH 7 > 6.99 @ 17.21°C 0755 Calibration completed. 0758 Marvin Saraccino arrives at gate. Elvin Chaven has not arrived yet. 6759 All depart to JPOP41S USOS 4000 arrived to JPOP415 approximately 12 cows are within 50 to 100 gds of JPOP415. 0806 Arrive at JPOPHIS. Waiting on

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MINE ESI, SR 279 LAGUNA PUBBLO,
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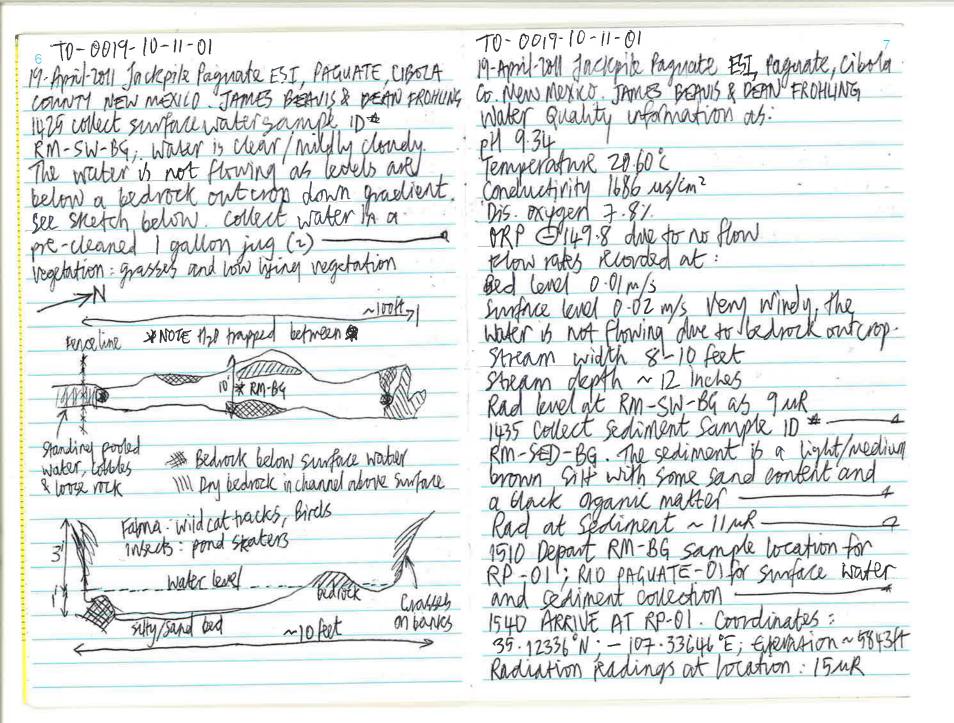
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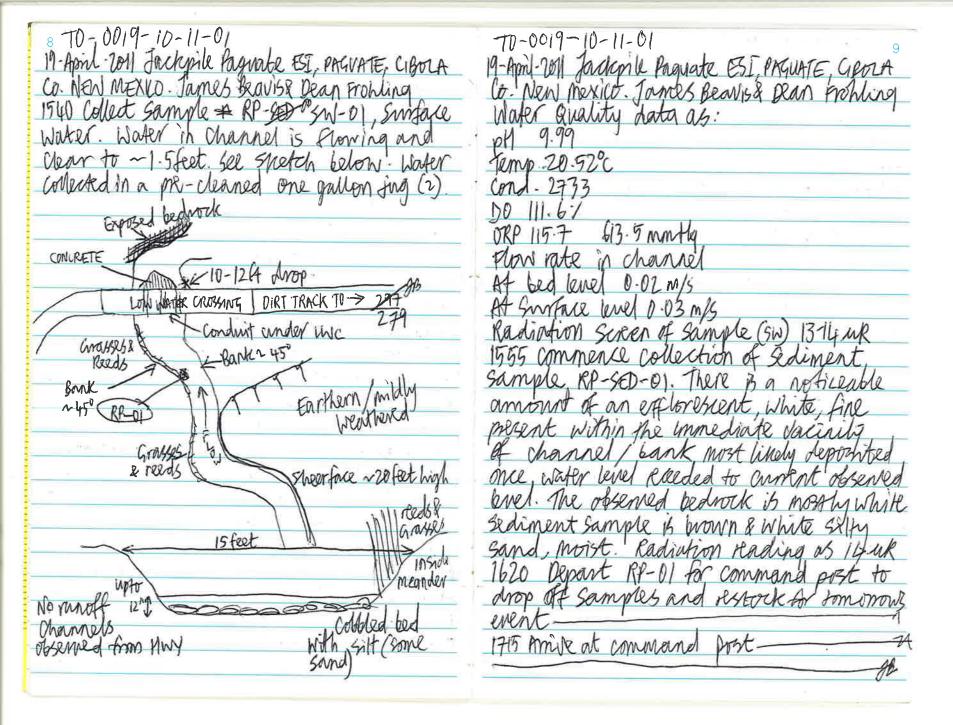
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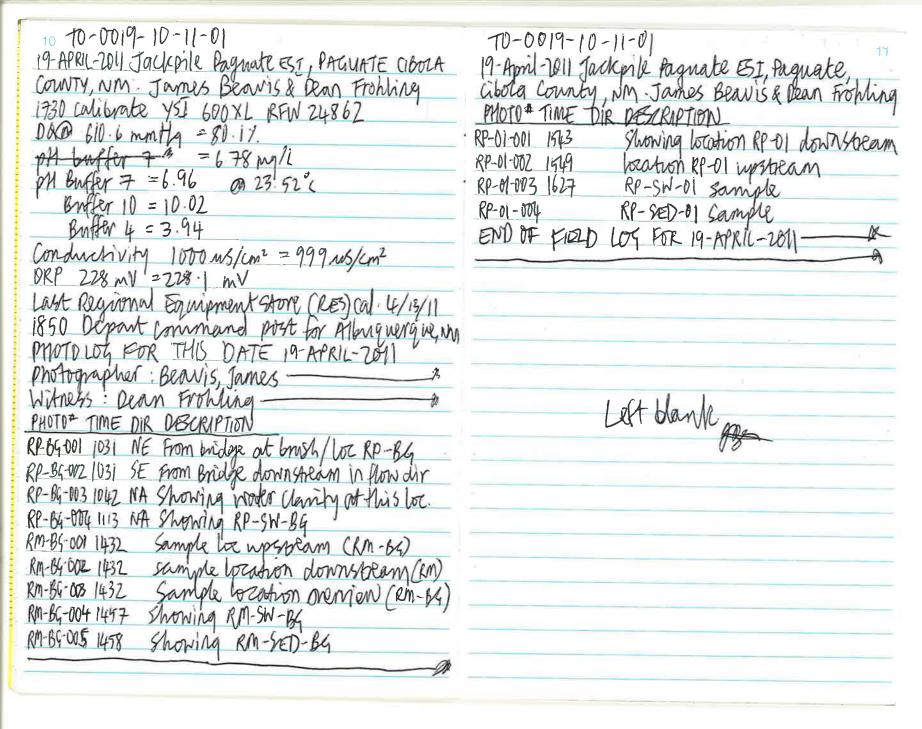
TDD# 0019-10-11-01 APRIL-19-2011 JACKPILE PAGNATE ESI, SR 279 LAGUNA PUEBLO, PAGUATE, CIBOLA COUNTY, NEW MEXICO SURFACE WATER AND SEDIMENT SAMPLING TEAM JAMES BEAVIS AND DEAN FROHLING. WEATHER (AM) ~59°F, MOSTLY SUNNY, WNW WIND ~20 mph (PM) HIGH OF 74°F, 10% CHANCE OF RAIN, WESTWIND 20-25 MPL 0700 START-3 BERNIS & PROHUNG Depart Moliday Inn Expert old town, Albuquerque, NM for the Josephile wanium mine 0800 Arrive at the Rublic Health Service, Indian health center (from here on referred to as command POST) and unitad sample media. Stock field Portal vehicle for sumpting event. EduiPMEN (: OGARMIN GPS map 76 CS EPA69774 2 SONY CHERSHOT 5.1 EPA 69942 Time/date-OK · LUDIUM MODEL 192 EPASGIZ45 · YSI Water quelity, 600XL RFW 24862 Jan-13-2012 · GED PUMP REWLOTS & FLOWINGER ID \$ 10816 0835 Field calibration YSI RPW 24862 Temp 18.61°C DQ: @ 611.8 mm Hg = 79.8%. 7.56mg/1 pH buffer 7, Rading 6.97 Buffer 4, reading 393 Buffer 10, Hading 9-47 Conductivity ORP: 288 mV, Rending 2-27.5 mV

TO-0019-10-11-01 19-April CON Jackepile Paswate ESI, Poguate, cibola county, NM. James Beavis & Dean Frohling 0925 PRF-sampling yealth & Safety discussion with PTL michelle from & the ground water Sampling team, Morpital is the Acoma service unit, ~28 miles on I-40, EXIT at Casino. Note the distance. Hazards include: working remote locations, snakes / scorprions etz. Radiation, USE SUNSCREEN, DO NOT INTERPER WITH CATTLE, FIRE HAZIFRISS INC. VEHICLE TAILPIPE ETC. MEUNTAIN LIONS AND OTHER LARGE ANIMALS, DO NOT DISTURB POTENTIAL TRIBAL ARTIFACTS ETC 0935 Background reading using Ludlum 192 at command post ~12 ul-0940 Depart command post for the RP-BG (RIO PAGUATE - Back ground) location with Anny Carcia from PABEO LÁGUNA ENVIRONMENT AL PROGRAM Amove at RP-BG coordinates of Sample Cocation 35.14095°N; -107.38322 € approx evelution 8264ft 1025 collect surface water sample RP-SW-BG worker is clear w/low problection pre-cleaned one gallon jud for filtering 0-65 micron high

TO-0019-10-11-01 TO-0019-10-11-01 19-April-2011 Jackpile Pagnate BI, Pagnate, 19-APRIL-2011 JACKPILE PAGUATE EST, PAGUATE, CIBOLA Citaly country, NM. James Bearis& Dean Frohling county, NEW MEXICO, James Beavis & DAN PROHLING Disposable geotech Gilter (73050004) using geostely geograph D& RFWUDIT. Collection Temperature 12.06°C regulation in wild 1025, april 485. court defails as Conductivity 818 MS/M2 & law lying veg Distribued orangen 71.7%. large trees, dudes ORP 115.1 & squinelly Collect sediment sample RP-SED-FG of Sand bank Mis wastion sediment is brown in color and fine / wedium crain size, Sandy / Loan Shallow/ ~ 15 Ret with high moisture content Stow (You precipitate/Efforvegrent ~30" Photos father out this location included a Courses day photolog Earthern bank 1125 surface water & sediment sampling ~294 team depart for command post for supplies mover level LACE ENTRY Rad reading of Sediment 7 uk/hr 1150 ARRIVEAT COMMAND POST 1230 LUNCH Rad @ lozatron ~ 12 WK 1315 Resumb operations, plan route to RM-BG. Rado NO 25 WR RIO MODUINO BACKGROUND. NOTE AM GARCIA WILL Stram width ~ 30 inchs NOT BE IN A ACLOMPANYING START-3 Mon rate at bed 0.03 m/s surface 0.25 m/s 1415 prine. coordinates of sample location 100 collect surface water screening Late 04: 35-15328°N, -107.39580E, approx elevation wring 15I water quality meter. Idta as: 5,970 St. Radiation radings @ sample overleaf. location ~ 10 mg/h



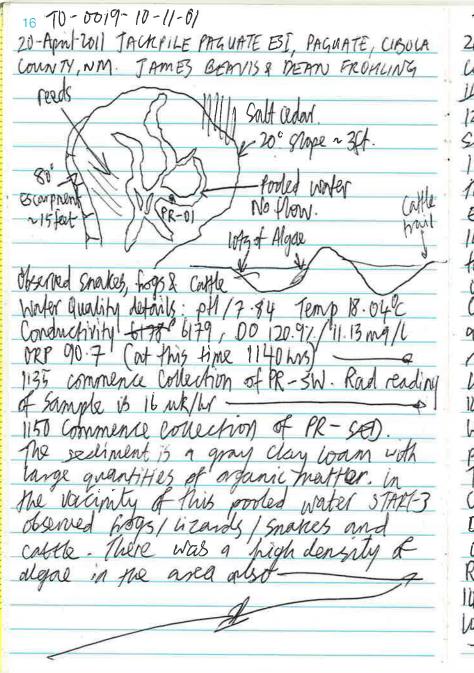




12 TDD# TO-0019-10-11-01 20-April 2811 Jackprile Pagnata ESI, PAGNATE, CIBOLA CO. NM Sirrface Water & sediment Sampling Jeam James Bearis & Dean Frohling (START-3). Weather Today: (Am) 50°F, SUNNY WITH WSW WINDS ~ 6mph (em) HILH ~ 77°F, MOSTLY SUNNY, SW WINDS ~ 15 mph 0700 STERT-3 Surface water and sediment Sampling team depart Holiday In Experts old town, Albuquerque, NM for Command posk, Vagung, NM-EDVIPMENT UST & PRESAMPLING CHECK : -· SONY CYBERSHOT 5.1 MPEG CAMERA EPA A69942 · Garmin GPS 76CS meloaded EPA A69774 · Ludlum model 192 Radmeter EPA S41245V - YSI 600 XL Water quality Meter RPW 24862V · GEDPUMP RFW21015V · Flow meter, 500 MATE 2000 10# 10816/ 0750 Pickup field supplies & check webside. 0800 Arrive at command post 0840 Morning START-3 health & Safety Ener awareness of surroundings, do not take chances when driving off road (pared) Fauna / flora, radiation - inextain. Fred in religious. In the field at each Weation discuss emergency plans w/

TO-0019-10-11-01 20-April-2011 Jackpule Paguate ESI, PAGUATE CIBOLACO. NM. James Beavist Dean Frohling 0855 Deport Command post for surface water and sediment sampling wation RP-07 (RIO PAGUATE -02) 0910 Arrive at RF-02 corondinates: 35.10304°N -107.32478°E, approximate elevation 5785ft. Byckground radiation reading 11-12MK/Lr The channel is dry with moist sediment in cattle footprints which are frequent phroughout the course of the channel in this Weation the bank/surfale is approximately 28-30" above the channel bed. See shotch below for defails: Radiution relorded as 15 mk at sample location Steep bank ~ 28" regetation: cultle footprints hnoughout Sout ledar. AN P Stoped bank ~ 25° RP-SED -02 o conthorn, grassed pedestals.

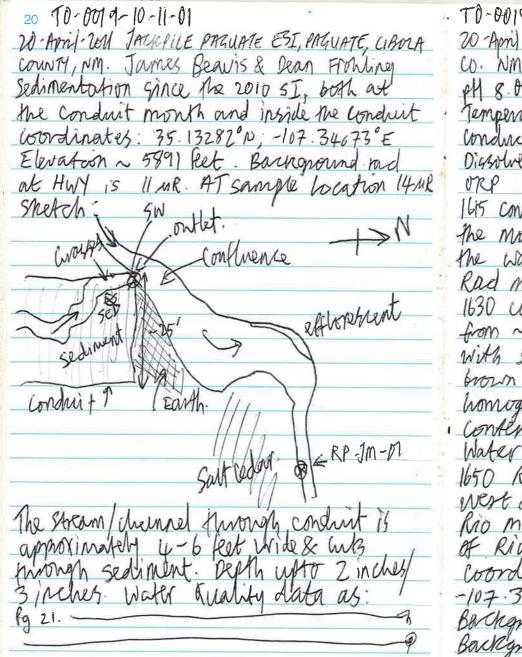
14 10-0019-10-11-01 TO-0019-10-11-01 20-April-2011 Jackpile Pagnate ESI, Pagnate, Cibola 20-April-2011 Jackpile Pagnate ESI, Pagnate, Gold County, NM James Bearis & Dear Frolling country, NM. James Bearis & Dean Frohling CRASS vanes 8-10ft Redogtal Cra45 & (harrel) Grasses & ted Salt RP-SED-02 salt cedar Cedar NO WATER A cattle thorofare 0425 Commence Collection of sediment sample RP-500-03 RP-SED-02. Rad reading of sample 13 mk 1948 Depart RP-02 for Surface water & sediment Sampling weather RP-03 goods of current, loca. Cattle fost prints throughout. Norre 1007 Aprine at RP-03. LATE ENTRY The toads observed within immediate vacinity. JESLA P Ment Sediment Collected as RP-SED-02 15 1 Solf cedar v150Ft. light brown silty/clay with minimal moisture Spurge -> Contentgregetations RP-500-03 Background reading at RP-03 recorded as: ~ 8 wk/hr. The channel is very dry. Dry channel Cattle posts Coordinates: 35.07877°N; -107.32771°E PACING NORTH Channel: 8-94 Wide, 2 approximate elevation, 5739 St. Rad 1035 Depart Sample Location RP-03 for readings, at channel as: 22 mg/m. PR-01 (PR=Pagnate Reservoir-01) See Sherch overleas. located sample over PR-01 coordinately DIO commence with collection ONR 35.07-299°N: -107.33237°E, 5740 ft elev. RP-SED-03. Rad reading of sample ~ 18 MR Background: 8 mg, lad Rading at PR-01:22 mg The sample is a Gray 18 rown sith day See shetch overleaf (pg. 16 for weation details. with Hitle/No moistne content.



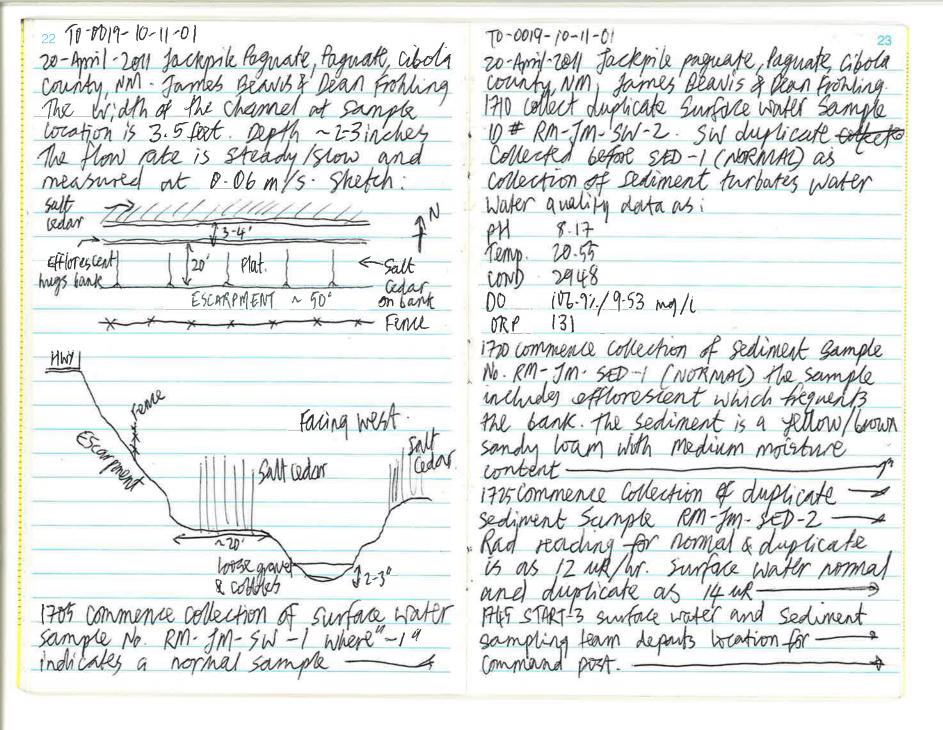
70-0019-10-11-01 20-APR-2011 Jacoppile Pagnate, Pagnate, cibola county, NM James Beauts & Dean Frohling 1400 Report pagnate Reservoir for Command part 1230 Aprive command post and unload Samples. Break for 'unch 1340 START-3 sampling team heads to The Merita diversion location. Follow EPA Labonna Tumer. 1400 Amine at the Mesita Diversion location for surface Water Sample collected under direction of EPA Turner. The Coordinates as: 35.02367°N; -107.32603°E approx devation: 5734 feet. Background sidiation 8 MR, Reading at Sample wation: 11 mg 1415 collect mesita Diversion surface Loyler sample 21.29°C Temp 656 10 4562 and. 114.6%, 10.05 mg/L DO ORP 1333 Rad reading of surface water lample: 7 mg 1425 Depart Mexita Diversion for sample Locations RM-JM, RP-JM & RP-JM-01

18 10-0019-10-11-01 20-April-2011 Jackpile Riguate BI, Pagnate, Cibola County, NM. James Beavis & Dean Frohling 1455 Amire at location. All three bocations and below a 45°, 50 food excarpment from The HWY 279 Bridge where RM is the RIO MOQUINO, RP is the RIO PAGUATE and RP-JM-01 falls after the confluence -1520 At Sediment and Surface water sample weation RP-JM-01 Que to moximity 5 TART-3 will collect the down shearn! location first. Coordinates: 35. 13280°N; -107.34650°E, Elevation ~5898 ft. Buildrenned raid reading on HWY 279 is 11 MR, at RP-JM-01 reading is 16 MR The channel is slowing at a medium velocity for an averview of location (smetch) see pg. 19 1540 commence collection of Simple water sample No. RP-JM-SW-01. Water Data Flow at channel bed: 0.16 m/s 0.15 m/5 Prow at surface: PM 8.11 Temp 20-49 Cond. 21.54 \$91./7.96 mg/L DO ORP 1297

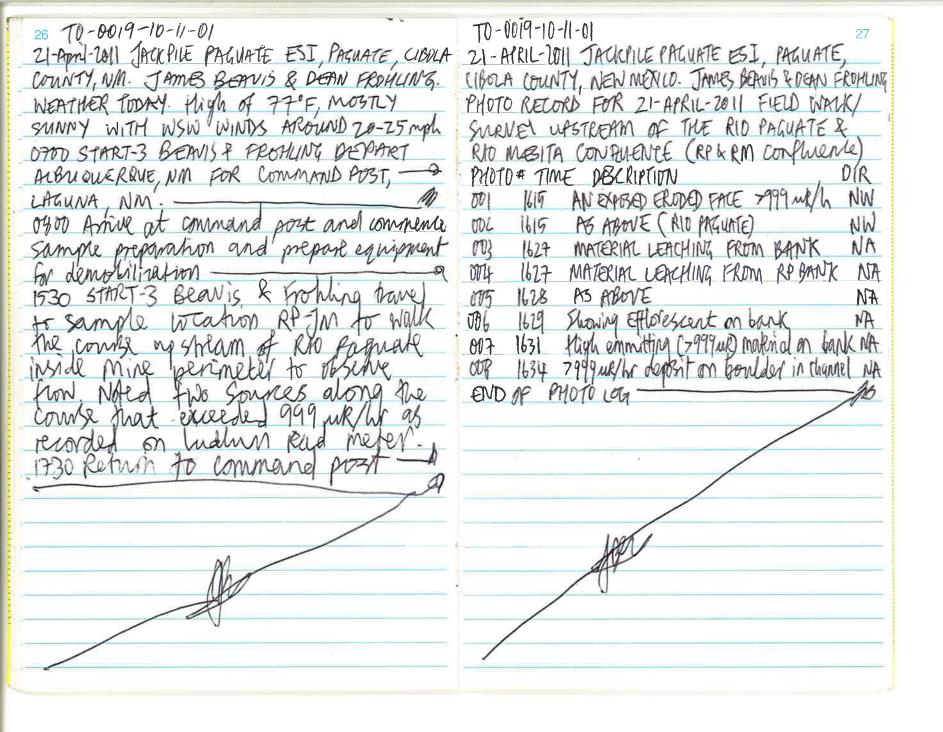
10-0019-10-11-01 20-April-2011 Jackpile Pagnate ESI, Pagnate Cibols county NM. James Boaris& Dean Fronting SWETCH ME AREA: · bank 2 Efflorescent Flow Conduit 10 m. deep. under road/bridge ~ 4 feet Grakes. Saltar Efflorestent 1555 Collect Sediment Cample RP-JM-500-01 the sediment is a yellow/ swown sandy loam with moisture content ~ moist. 605 Relocate approximately 25 feet west to the Riv laguate (RPJIM) sampling location. The Water is flowing through a Conduit Cr25 feet diameter). The bocation is observed to have considerable



- TO-0019-10-11-01 20 April 2011 JACKPILE PAGUATE ESI, PAGUATE, CIBOLA CO. NM. James Bearis & Dean Frohling PH 8.09 Temperature 17-89°C Conductivity 1336 Dissolved Pxy 82.5/ /7.81 mg/1 orp 128 6 1615 commence collection of RP-JM-SW from The month / lip of the conduit before the water enters the cherrel/confluence. Rad meter reading of SW is 11 up/hr 1630 collect sediment sample RP-JM-SED from ~ 4 feet inside conduit at interface with stream flow. The sediment is a brown Sadiment with high sand content, homogenous shroughout. Ligh moisture Content Water flow is at 0. 73 m/second. 1650 Relocate approximately 150-200 feet west of RP-JM" upstream on the Kio moguino before the confluence of Rio Mognins & Rio Pagnaft Coordinates are: 35. 13318 N and -107.34701°E. Dev: 5891 St. Background and at MWY is HUR Background at Sample location is Iluk



10-0079-10-11-01 70-0019-10-11-01 20 April-2011 Jackpile Pagnate Est, fagnate, Cibola Conney, NM. James Bearis & Dean Frohing 20-April-2011 JACKPILE PAGUATE ESI, PAGUATE, CIBOLA LOUNTY, NM. James Bearis & Dean Frohing 1800 ARRIVE AT COMMAND POST AND UNLOPED PHOTO 10# TIME DIR DESCRIPTION 4AMPLES, DECON EQUIPMENT FOR DEMOSE PR-08 1158 NA Showing PR-SED& PR-SW 1850 Report ammand post for Albuquerque Moirersim-of 1403 overview at the Messita Diversion approx one how travel time. ADjust Sion-OL 1404 A3 above 2000 END DAY .-Moirersion of 148 NA Megita diversion suntence water PHOTOLOG FOR 20-APRIL-2011 RP-JM-11-01 1550 SE Facing Downstream at RP-JM-01 PHOTOGRAPHER: James BEAVIS RP-JM-01-02 1550 NW Facing RP&RM Confluence WITNESS: DEAN FROHLING RP-JM-01-03 1550 NA Overview of RP-JM-01 sample location PHOTO ID (tentative) SITE ID - No. sequence RP-JM-OTOY ILL NA RP-JM-71-SW PHOTO ID TIME DIR DESCRIPTION RP-JM-0-05 1612 NA RP-JM-01-50 RP-02-01 1932 S Facing down stream of RPOZ A RP-JM-09001 1612 S Rio Enquale through conduit RP-02-02 0934 N Facing uposteam at RFD2 RP-7M-02 1612 NA RP Flow out/at mouth of Conduit RP-02-03 8936 NA RP-02-SED RP-JM-03 1613 Chancel & sidiment of RP 5 RP-03-01 1018 S Facing obown stram at RP03 RP-JM-04 1616 SW SIMPLY-3 OSURETING SW Sample RP-03-02 1078 Facing upspeam at RPD3 RP-JM-05 1633 NA RP-JM-SW & RP-JM-SED RP-03-03 1023 NA RP-03-50 RM-JM-01 1657 E START-3 measuring RM flowlate RP-03-04 1025 E Overview of semple preation RM-1m-02 1658 E dannistram fring confluence PR-01 1657 NA WILDLIFE at pasenvoir - SWARE RM-JM-03 1658 w facing upstream PR-02 1138 MA Wildlife of Rosenjoir - Frog RM-JM-04 1734 NA RM-JA-5W-1 (Normal) PR-03 1138 PVSW Pooled With within Reservoir bank. RM-JM-0\$ 1737 NA RM-JM-SED-1 (Normal) PR- 04 1/35 N Overview of Sermpling Lozation RM-JM-02 1737 NA RM-JM-SW-2 (Augheate) PR- 175 1199 MA Wildlife at Reservoir 2nd snake Rm-Jm-07 1740 NA Rm-Jm-SW-2 (duplicate) PR- OR 1170 NW DUENTIEW of WEEKEN PR END OF PHOTOGOG PR-17 1150 NA Different Vorniety of Snake



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QUALITY ASSURANCE SAMPLING PLAN

FOR

EXPANDED SITE INSPECTION

JACKPILE-PAGUATE URANIUM MINE SR 279 LAGUNA PUEBLO PAGUATE, CIBOLA COUNTY, NEW MEXICO

Prepared for

U.S. Environmental Protection Agency Region 6

Linda Carter, Project Officer 1445 Ross Avenue Dallas, Texas 75202

Contract No. EP-W-06-042
Technical Direction Document No. TO-0019-10-11-01
WESTON Work Order No. 20406.012.019.0603.01
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FPN No. N/A
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March 2011

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1. INTRODUCTION

Weston Solutions, Inc. (WESTON®), the EPA Region 6 Superfund Technical Assessment and Response Team (START-3) contractor, has been tasked by the U.S. Environmental Protection Agency (EPA) Region 6 Prevention and Response Branch (PRB) under Contract No. EP-W-06-042 and Technical Direction Document (TDD) No. TO-0019-10-11-1 (Appendix E) to perform Expanded Site Investigation (ESI) activities at the Jackpile-Paguate Uranium Mine site located on SR 279 Laguna Pueblo in Paguate, Cibola County, New Mexico. A Site Location Map is provided as Figure 1-1. The Comprehensive Environmental Response, Compensation, and Liability Information System (CERCLIS) database number assigned to the Jackpile-Paguate Uranium Mine Site is NMN000607033. START-3 has prepared this Quality Assurance Sampling Plan (QASP) to describe the technical scope of work (SOW) to be completed as part of the ESI.

1.1 PROJECT OBJECTIVES

START-3 is providing technical assistance to EPA Region 6 for the performance of an ESI at the Jackpile-Paguate Uranium Mine site. The objective of the ESI is to identify potential threats that hazardous substances attributable to the site may pose to human health and the environment. START-3 will assess the existence and migration of hazardous substances and identify the receptors, or targets, potentially exposed to the hazardous substances.

The ESI objective will be achieved by evaluating data obtained during the 2010 Site Inspection (SI), along with new analytical data from surface water and groundwater samples collected at onsite, downgradient, and background locations. Sediment samples in the surface water pathway may also be collected during this ESI. The site-specific Contaminants of Concern (CoCs) include isotopic uranium, total metals (including mercury), gross alpha, beta, and gamma, and radium 226 + 228. The site-specific screening levels will be identified as three times the maximum background sample concentrations.

In addition, water samples will be analyzed for total dissolved solids and general cation/anion chemistry. These data will be used to evaluate sample locations with a conceptual site model that considers hydrologic and hydrochemical data.

The information collected by START-3 during the ESI will be used to evaluate the site using the Hazard Ranking System (HRS) and to assist in determining whether the site is a potential candidate for inclusion on the National Priorities List (NPL). The intent of the ESI is to provide documentation necessary to either rank the site on the NPL or assign a "No Further Remedial Action Planned" (NFRAP) status.

1.2 PROJECT TEAM

The Project Team will consist of START-3 personnel including Michelle Brown as the Project Team Leader (PTL), a Field Safety Officer (FSO), a Data Manager, and additional START-3 Field Team Members to assist with sampling activities. The PTL will oversee collection of the samples as necessary, record the activities at each sample location in the field logbook, and verify sample documentation. Sample documentation and preparation is also the responsibility of START-3. The PTL will be responsible for documenting the work performed and will serve as START-3 liaison to EPA Region 6.

1.3 QASP FORMAT

This QASP has been organized in a format that is intended to facilitate and effectively meet the objective of the removal assessment. The QASP is organized as follows:

- Section 1 Introduction
- Section 2 Site Background
- Section 3 Sampling Approach and Procedures
- Section 4 Analytical Methods
- Section 5 Data Validation
- Section 6 Quality Assurance

Tables are included at the end of each representative section. All figures are provided as separate Portable Document Format (PDF) files. Appendices are attached with the following information:

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Appendix A EPA Contract Laboratory Program Guidance for Field Samplers

Appendix B Site-Specific Data Quality Objectives

Appendix C WESTON and ERT Standard Operating Procedures

Appendix D Laboratory Analyte List and Reporting Limits

Appendix E START-3 TDD No. TO-0019-10-11-1

2. SITE BACKGROUND

Information regarding the site location, description, and site history are included in the following

subsections.

2.1 SITE LOCATION AND DESCRIPTION

The Jackpile-Paguate Uranium Mine site consists of approximately 7,868 acres of uranium mine

leases within the Pueblo of Laguna, Paguate, Cibola County, New Mexico. The site is located 40

miles west of Albuquerque, New Mexico and contains 2,656 acres of disturbed land that consists

of three open pits and nine underground mines. The geographical center of the site is Latitude

35°8'16.74" North and Longitude 107°20'51.84" West. A Site Area Map derived from the ESRI

geographical information system (GIS) Microsoft Virtual Earth Roads, 2009 is included as

Figure 2-1.

The village of Paguate is located adjacent and west of the site with a population of approximately

474 according to the 2000 U.S. Census. The remaining surrounding area is mainly undeveloped,

with the perennial rivers Rio Moquino and Rio Paguate bisecting the site. The Rio Paguate

continues south of the former uranium mine continuing through the Paguate Reservoir, 4 miles

south of the site, into the Rio San Jose.

2.2 SITE HISTORY

The mine was operated by Anaconda Minerals Company, a division of the Atlantic Richfield

Company, from 1953 through 31 March 1982. The mine was closed because of a sustained fall in

the price and demand for uranium.

During its operation, open pit mining was conducted with large front-end loaders and haul

trucks. The overburden, consisting of topsoil, alluvium, shale, and sandstone, was blasted or

ripped, removed from the open pits, and placed in waste dumps. Underground mining was

conducted by driving adits, or declines, to the ore zone. The ore was removed by modified room

and pillar methods. Mine water was collected in sumps and pumped to ponds in the open pits.

Waste rock was placed in waste dumps. The ore was segregated and shipped by rail to the

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TDD NO. TO-0019-10-11-01

JACKPILE-PAGUATE QASP.DOC

2-1

Anaconda mill, 40 miles west of the mine. Ore was also trucked out in open top trainers which were supposed to be covered but most of the times were not. Therefore, ore was scattered along old highway 279 from the numerous accidents on the hill called Questa.

During the 29 years of mining, approximately 400 million tons of earth was moved within the mine area, and about 25 million tons of ore was removed.

3. SAMPLING APPROACH AND PROCEDURES

The specific field investigation activities that will be conducted during site sampling are presented in the following subsections. Samples collected as part of this ESI will be obtained in accordance with EPA *Contract Laboratory Program Guidance for Field Samplers*, *EPA540-R-00-003*, *OSWER 9240.0-35* (Appendix A).

3.1 OVERVIEW OF MONITORING/SAMPLING ACTIVITIES

This ESI is being performed to evaluate the surface water and groundwater-to-surface water exposure pathways at or near the Jackpile-Paguate Uranium Mine site. The primary CoCs at the site are uranium, gross alpha, beta, and gamma (which includes thorium), radium 226+228, and heavy metals. Subcontracted laboratories will be utilized for analysis of isotopic uranium; total metals (including mercury); gross alpha, beta, and gamma; and radium 226+228.

In addition, water samples will be analyzed for total dissolved solids (TDS) and general cation/anion chemistry. These data will be used to evaluate the samples in the context of the site hydrologic model.

The EPA SAM and START-3 will collect 11 (including duplicates) surface water samples from Rios Paguate and Moquino, and 14 (including duplicates) groundwater samples from existing wells around the site. START-3 will also collect ten sediment samples from the surface water pathways that include the Rio Moquino, Rio Paguate, Paguate Reservoir, and the Rio San Jose (including background and Quality Assurance/Quality Control [QA/QC]). If necessary, sampling of investigation-derived waste (IDW) will be performed to properly dispose of IDW. START-3 will use EPA Scribe Environmental Sampling Data Management System (SCRIBE) software to manage sample data.

3.1.1 Data Quality Objectives

The objectives of the sampling activities described in this QASP are to collect representative surface water and groundwater samples to determine if there has been a release of site-related hazardous substances and contaminants into the surface water and groundwater pathways. To

accomplish this, the following data quality objectives (DQOs) have been established and are included in Appendix B:

- Determine if there is a release of site-related contamination in the groundwater pathway.
- Determine if there is a release of site-related contamination in the surface water pathway.
- Provide water chemistry data to support hydrologic model.

The DQOs were developed using the seven-step process set out in the *EPA Guidance for Quality Assurance Project Plans: EPA QA/G-5*. Table 3-1 summarizes the sample locations and the rational for their collection.

3.1.2 Health and Safety Plan Implementation

The START-3 field activities will be conducted in accordance with the site-specific health and safety plan (HASP). The FSO will be responsible for implementation of the HASP during all field investigation activities. In accordance with the WESTON general health and safety operating procedures, the field team will also drive the route to the hospital specified in the HASP prior to initiating sampling activities. Radiation monitoring will be conducted at all sample locations prior to and during sample collection using a MicroR meter.

3.1.3 Tribal Involvement

The EPA SAM will coordinate all assessment activities with the Pueblo of Laguna Environment Department. Any issues will be directed to the EPA SAM. If the EPA SAM is not present, the START-3 PTL, under the guidance of the START-3 Assessment/Inspection Manager, will manage tribal involvement in the field as directed by the SAM. If a tribal involvement plan and implementation program becomes necessary, START-3 will establish each if requested by the EPA SAM. START-3 will work as directed by the EPA SAM to obtain access to all designated sampling locations.

3.2 SAMPLING/MONITORING APPROACH

All samples will be collected in general accordance with the WESTON Standard Operating Procedures (SOPs) 1002-01 for Surface Water Sample Collection, 1002-02 for Groundwater

Monitoring Well Sample Collection, and 1002-04 for Sediment Sampling (Appendix C). The specific sampling procedures are described below.

3.2.1 Groundwater Pathway Sampling

Fourteen groundwater samples (including two duplicates) will be collected from existing monitoring wells. START-3 will measure depth to groundwater in each of the wells and then follow the EPA *Guidance for Low-Flow* (Minimal Drawdown) *Groundwater Sampling Procedures* (Appendix D) for sampling the wells, if appropriate. START-3 anticipates using either a Smeal rig, QED pneumatic pump or peristaltic pump depending on the depth to groundwater. Readings for temperature, pH, and conductivity will be collected every 5 minutes. Once three consecutive readings stabilize for pH (± 0.5 units), conductivity (± 10% µmhos/cm), and temperature (± 1°C), or the water has purged for a maximum of 30 minutes, the samples will be collected.

The exact number of samples and locations of the samples will be decided by the EPA SAM and START-3 PTL. The proposed groundwater sample locations are presented in Figure 3-1.

The groundwater samples will be submitted to a National Environmental Laboratory Accreditation Program (NELAP) certified laboratory, for the following analyses:

- Isotopic Uranium (including isotopes 234, 235, and 238) by ASTM 3972-90M alpha spectrometry.
- Total metals including mercury by SW846 Methods 6010B and 7470/7471.
- Gross alpha, beta, and gamma (which include thorium) radiation by EPA Method 9310.
- Radium 226 and 228 by EPA Method 903.1.
- Total Dissolved Solids (TDS) by EPA Method 160.1
- General Anion Chemistry by EPA Method 9056. The laboratory will also calculate total alkalinity and carbonate and bicarbonate calculations.

Laboratory-specific analyte lists and reporting limits are included in Appendix D. Deviations from the sample locations may occur at the EPA SAM's direction, due to new observations made prior to sampling, information obtained in the field that warrants an altered sampling point,

difficulty in sample collection, or limited access. The EPA SAM will be notified, and concurrence will be obtained should significant deviations from the planned sampling points be proposed. Details regarding deviations of the QASP will be documented in the site logbook.

3.2.2 Surface Water Pathway Sampling

START-3 will collect 11 surface water, and 10 sediment samples (including two background and one duplicate sample) as part of the ESI to document a release to the surface water pathway from the site. The proposed surface water and/or sediment sample locations are presented in Figure 3-1.

Surface water and sediment samples will be submitted to a NELAP certified laboratory to be determined for the following analyses:

- Isotopic Uranium (including isotopes 234, 235, and 238) by ASTM 3972-90M alpha spectrometry.
- Total metals including mercury by SW846 Methods 6010B and 7470/7471, respectively.
- Gross alpha, beta, and gamma (including thorium) radiation by EPA Method 9310.
- Radium 226 +228 by EPA Method 903.1.

In addition, surface water samples will be analyzed for the following:

- Total Dissolved Solids (TDS) by EPA Method 160.1
- General Anion Chemistry by EPA Method 9056. The laboratory will also calculate total alkalinity and carbonate and bicarbonate calculations.

The laboratory-specific analyte list and reporting limits are included in Appendix D.

3.2.3 Investigation-Derived Wastes

Attempts will be made to eliminate or minimize generation of IDW during this investigation. All non-dedicated equipment will be decontaminated according to WESTON SOP 1201.01. Non-dedicated equipment will be rinsed with soap and water and attempts will be made to dispose of decontamination fluids and purge waters on-site. The analytical data from collected samples will be reviewed after completion of the field activities, and disposal options will be evaluated

accordingly. It is anticipated that minimal amounts of IDW will be generated during this activity.

3.2.4 Sampling and Sample Handling Procedures

Samples will be collected using equipment and procedures appropriate to the matrix, parameters, and sampling objectives. All water samples collected will be field filtered using 0.45µ filter paper according to EPA Emergency Response Team (ERT) SOP #2007 (Appendix C). The volume of the sample collected must be sufficient to perform the laboratory analysis requested. Samples must be stored in the proper types of containers and preserved in a manner appropriate to the analysis to be performed. A sample collection and analyses summary table is presented in Section 4.

All clean, decontaminated sampling equipment and sample containers will be maintained in a clean, segregated area. All samples will be collected with clean decontaminated equipment following WESTON SOP 1201.01. All samples collected for laboratory analysis will be placed directly into pre-cleaned, unused glass or plastic containers. Sampling personnel will change gloves between each sample collection/handling. All samples will be assembled and catalogued prior to shipping to the designated laboratory (following WESTON SOP 1101.1 and 1102.01).

3.2.5 Quality Assurance/Quality Control Samples

START-3 will collect field duplicate and Matrix Spike/Matrix Spike Duplicate (MS/MSD) samples of groundwater, surface water, and solid-matrix samples, as well as prepare equipment rinsate blank samples as needed during the assessment sampling activities. QA/QC samples will be collected according to the following dictates:

- Blind field duplicate samples will be collected during sampling activities at locations selected by the START-3 PTL. The data obtained from these samples will be used to assist in the quality assurance of the sampling procedures and laboratory analytical data by allowing an evaluation of reproducibility of results. Efforts will be made to collect duplicate samples in locations where there is visual evidence of contamination or where contamination is suspected. Blind field duplicate samples will be collected at the rate of one duplicate for every 10 samples collected.
- Equipment rinsate blanks will be prepared by pouring laboratory-grade deionized water over nondisposable sampling equipment after it has been decontaminated and collecting the rinse

water in sample containers for analyses. These samples will be prepared to demonstrate that the equipment decontamination procedures for the sampling equipment were performed effectively. The equipment rinsate blanks will be prepared each day that non-disposable sampling equipment is used.

- Temperature blanks will be prepared in the field and will consist of one 40-milliliter glass sample container with Teflon-lined septum cap. The temperature blank will be packaged along with the field samples in the shipping cooler and will represent the temperature of the incoming cooler upon receipt at the laboratory. Use of these samples within a shipping container enables the laboratory to assess the temperature of the shipment without disturbing any of the field samples.
- MS/MSD samples will be collected during sampling activities at locations selected by the START-3 PTL. The data obtained from these samples will be used to assist in the quality assurance of the sampling procedures and laboratory analytical data by allowing an evaluation of reproducibility of results. Efforts will be made to collect MS/MSD samples in locations where there is no visual evidence of contamination or where contamination is not suspected. MS/MSD samples will be collected at the rate of one MS/MSD sample per matrix for every 20 samples collected.

3.3 SAMPLE MANAGEMENT

Specific nomenclature that will be used by START-3 will provide a consistent means of facilitating the sampling and overall data management for the project (WESTON SOP 0110.05). The START-3 Assessment/Inspection Manager must approve any deviations from the sample nomenclature proposed below.

As stated in WESTON SOP 0110.05, sample nomenclature will follow a general format regardless of the type or location of the sample collected. The general nomenclature consists of the following components:

- Geographic location (e.g., location within a surface water body or monitoring well).
- Collection type (composite, grab, etc.).
- QA/QC type (normal, duplicate, etc.).
- Sequence An additional parameter used to further differentiate samples.

Sample data management will be completed utilizing SCRIBE including Chain-of-Custody (COC) and sample documentation needs.

3.4 DECONTAMINATION

The nondisposable sampling equipment (hand trowels, stainless steel bowls, etc.) used during the sample collection process will be thoroughly decontaminated before initial use, between use, and at the end of the field investigation. Equipment decontamination will be completed in the following steps:

- Water spray or brush, if needed, to remove soil/sediment from the equipment.
- Nonphosphate detergent and potable water wash to clean the equipment.
- Final potable water rinse.
- Air-dried equipment.

Personnel decontamination procedures are described in the site-specific HASP. All decontamination activities will be conducted at a temporary decontamination pad that will be constructed in an area to be determined by the PTL prior to the beginning of field activities.

Excess soil and fluids generated as a result of equipment decontamination will be placed in a drum and staged in an area to be determined by the PTL. The drum will be labelled on the side with the name of the site, the contents, sampling location, and date.

3.5 SAMPLE PRESERVATION, CONTAINERS, AND HOLD TIMES

Once collected, samples will be stored in coolers on-site until shipped for laboratory analysis. Samples designated for alpha spectrometry analysis do not require maintenance of a specific temperature range and do not have a holding time limit. Chain-of-custody forms will be completed for each sample shipment and sent with the samples to the designated laboratory by overnight carrier.

START-3 will receive analytical results based on discussions with the EPA SAM. This turnaround time (TAT) is initiated when the samples are received by the laboratory and continues until the analytical results are made available to START-3 either verbally or by providing facsimile or email copies of the results for review. Samples that have been analyzed will be disposed by the designated laboratory in accordance with the laboratory SOPs.

Table 3-1 Sample Locations and Sampling Rationale Jackpile-Paguate Uranium Mine Paguate, Cibola County, New Mexico

Sample Name	Sample Matrix	Sample Location (refer to Figure 3-1)	Rationale
RM-SW-BG-(date) RM-SED-BG-(date)	Surface water Sediment	Rio Moquino upstream of Jackpile Mine	Previous sample location. Collected to establish background concentration on Rio Maoquino.
RP-SW-BG-(date) RP-SED-BG-(date)	Surface water Sediment	Rio Paguate upstream of Jackpile Mine	Previous sample location. Collected to establish background concentration on Rio Maoquino.
RM-JM-SW-(date) RM-JM-SED-(date)	Surface water Sediment	Rio Moquino within the mine boundaries, upstream of its confluence with Rio Paguate	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water
RP-JM-SW-(date) RP-JM-SED-(date)	Surface water Sediment	Rio Paguate within the mine boundaries, upstream of its confluence with Rio Moquino	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water
RP-JM-SW-01-(date) RP-JM-SED-01	Surface water Sediment	Rio Raguate after the confluence of it with Rio Moquino and within mine boundaries.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water
RP-SW-01-(date) RP-SED-01-(date)	Surface water Sediment	Rio Paguate on the southern end of the mine boundary	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water
RP-SW-01-(date) RP-SED-01-(date)	Surface water Sediment	Rio Paguate on the southern end of the mine boundary	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water
RP-SW-02-(date) RP-SED-02-(date)	Surface water Sediment	Rio Paguate downstream of Jackpile Mine.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water
RP-SW-02-(date) RP-SED-02-(date)	Surface water Sediment	Rio Paguate downstream of Jackpile Mine.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water
PR-SW-01-(date) PR-SED-01-(date)	Surface water Sediment	Paguate Reservoir (aka Mesita Reservoir) downstream of Jackpile Mine.	Previous sample location. This sample location established an observed release during 2010 SI sampling. Collected to establish migration of contaminants from mine to surface water
MR-SW-(date)	Surface water	Irrigation canal off the Mesita Reservoir (aka Paguate Reservoir) downstream of Jackpile Mine. (not on figure – to be determined in field)	This sample location is being collected per request from the Village of Mesita
MW-RM-(date)	Groundwater	Monitoring well located off Rio Moquino, upgradient of Jackpile Mine.	Collected to establish background concentration upgradient and north of Jackpile Mine.
MW-1-(date)	Groundwater	Monitoring well located upgradient and to the northwest of Jackpile Mine.	Collected to establish background concentration upgradient of Jackpile Mine.
MW-8-(date)	Groundwater	Monitoring well located upgradient and to the west of Jackpile Mine.	Collected to establish background concentration upgradient of Jackpile Mine.

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Sample Name	Sample Matrix	Sample Location (refer to Figure 3-1)	Rationale
MW-7-(date)	Groundwater	Monitoring well located upgradient and to the west of Jackpile Mine.	Collected to establish background concentration upgradient of Jackpile Mine.
SPOP35-(date)	Groundwater	Monitoring well located within South Paguate Pit	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations
Well#4-(date)	Groundwater	Monitoring well located within Jackpile Mine along Rio Paguate	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations
MW-3-(date)	Groundwater	Monitoring well located within Jackpile Mine along Rio Paguate	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations
NPOP20E-(date)	Groundwater	Monitoring well located within North Paguate Pit	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations
JPOP41N-(date)	Groundwater	Monitoring well located within Jackpile Pit	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations
MW-2-(date)	Groundwater	Monitoring well located just south and downgradient of mine	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations
MW-6-(date)	Groundwater	Monitoring well located south and downgradient of mine	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations
MW-MD-(date)	Groundwater	Monitoring well located downgradient of mine and west of Mesita Dam	Collected to determine whether CoCs identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations

4. ANALYTICAL METHODS

START-3 will collect samples for both radiological and chemical analysis. Tables 4-1 and 4-2 summarize the samples that will be collected, including the volumes, container types, and associated analytical methods.

4.1 RADIOLOGICAL ANALYSIS

Samples collected by START-3 during the ESI will be analyzed by a NELAP certified laboratory. Surface water, sediment, and groundwater samples will be submitted for the following analyses:

- Isotopic Uranium (including isotopes 234, 235, and 238) by ASTM 3972-90M alpha spectrometry (groundwater only).
- Gross alpha, beta, and gamma radiation by EPA Method 9310.
- Radium 226 + 228 by EPA Method 903.1.

4.2 CHEMICAL ANALYSIS

Samples collected by START-3 during the ESI will be analyzed by NELAP certified laboratory. Surface water, sediment, and groundwater samples will be submitted for the following analyses:

- Total Metals by SW-846 Method 6010B
- Mercury by SW-846 Method 7470/7471.

In addition, surface water and groundwater will be submitted for the following analyses:

- Total Dissolved Solids (TDS) by EPA Method 160.1
- General Anion Chemistry by EPA Method 9056.

Table 4-1 Requirements for Containers, Preservation Techniques, Sample Volumes, and Holding Times Jackpile-Paguate Uranium Mine Paguate, Cibola County, New Mexico

Name	Analytical Methods	Container	Preservation	Minimum Sample Volume or Weight	Maximum Holding Time
TAL Metals and Mercury	SW846 6010B and SW846 7470/7471	Polyethylene (water), Glass (solid)	HNO ₃ to pH<2 (water), 4°C	500 mL, 80z	28 days for mercury 180 days all other metals
General Chemistry (anions)	SW846 9056	Polyethylene (water)	NA	500 mL,	28 days
Total Dissolved Solids	SW846 160.1	Polyethylene (water)	NA	500 mL	7 days
Uranium (isotopic)	Alpha Spec ASTM 3972-90M	Polyethylene (water), Glass (solid)	HNO ₃ to pH< 2 (water), NA (soil)	1 liter, 8 oz	6 months
Gross alpha, beta, gamma	SW-846 9310	Polyethylene (water), Glass (solid)	HNO ₃ to pH< 2 (water), NA (soil)	1 liter, 8 oz	NA
Radium 226+228	SW-846 903/9315	Polyethylene (water), Glass (solid)	HNO ₃ to pH< 2 (water), NA (soil)	1 liter, 8 oz	NA

Table 4-2 Field and Laboratory QA/QC and Analysis Summary Jackpile-Paguate Uranium Mine Paguate, Cibola County, New Mexico

Comple	Sample Collection	No. of		
Sample Type	Sample Collection Method	No. 01 Samples	Rationale	EPA Analytical Method
Groundwater	SOP 1002.2	8	Extent of contamination	SW846 6010B SW846 7470/7471 SW846 7199 ASTM 3972-90M SW846 9310 SW846 903 SW-846 160.1
Groundwater	SOP 1002.2	4	Background	SW846 6010B SW846 7470/7471 SW846 7199 ASTM 3972-90M SW846 9310 SW846 903 SW-846 160.1
Duplicate Groundwater (10%)	SOP 1002.2	2	QA/QC	SW846 6010B SW846 7470/7471 SW846 7199 ASTM 3972-90M SW846 9310 SW846 903 SW-846 160.1
Surface Water (water)	SOP 1002.1	7	Extent of contamination	SW846 6010B SW846 7470/7471 SW846 7199 SW846 9310 SW846 903 SW-846 160.1
Surface Water (water)	SOP 1002.1	1	Village of Mesita request	SW846 6010B SW846 7470/7471 SW846 7199 SW846 9310 SW846 903 SW-846 160.1
Surface Water (water)	SOP 1002.1	2	Background	SW846 6010B SW846 7470/7471 SW846 7199 SW846 9310 SW846 903 SW-846 160.1
Duplicate Surface Water (10%)	SOP 1002.1	1	QA/QC	SW846 6010B SW846 7470/7471 SW846 7199 SW846 9310 SW846 903 SW-846 160.1
Surface Water (sediment)	SOP 1002.4	7	Extent of contamination	SW846 6010B SW846 7470/7471 SW846 9310 SW846 903
Surface Water (sediment)	SOP 1002.4	2	Background	SW846 9310 SW846 903
Duplicate Sediment (10%)	SOP 1002.4	1	QA/QC	SW846 6010B SW846 7470/7471 SW846 9310 SW846 903
Rinsate Blanks ¹	N/A	5	QA/QC	SW846 6010B SW846 7470/7471 SW846 9310 SW846 903

Table 4-2 (Continued) Field and Laboratory QA/QC and Analysis Summary Jackpile-Paguate Uranium Mine Paguate, Cibola County, New Mexico

Sample Type	Sample Collection Method	No. of Samples	Rationale	EPA Analytical Method
Temperature Blanks ²	N/A	5	QA/QC	Vial temperature measured upon arrival at laboratory

Note:

¹Rinsate blanks: samples will be collected at the rate of 1 per day per nondisposable sampling equipment.

²Temperature blanks: samples will be collected at a rate of 1 per cooler.

5. DATA VALIDATION

START-3 will validate the radioanalytical data by having each data set reviewed by a professional health physicist. A summary of the data validation and findings will be presented in Summary Reports as part of the final report. START-3 will evaluate the following to verify that the radioanalytical data are within acceptable QA/QC tolerances:

- The completeness of the Laboratory Reports, verifying that all required components of the report are present and that the samples indicated on the accompanying chain-of-custody are addressed in the report.
- The results of laboratory blank analyses.
- The results of laboratory control sample (LCS) analyses.
- Compound identification and quantification accuracy relative to expected isotopic ratios for uranium and its decay products.
- Laboratory precision, through review of the results for blind field duplicates.

The inorganic analytical data generated by the designated laboratory will be validated using EPA-approved data validation procedures in accordance with the EPA *CLP National Functional Guidelines* for Inorganic Data Review (October 2004). A summary of the data validation findings will be presented in Data Validation Summary Reports as part of the final report. The following will be evaluated to verify that the analytical data is within acceptable QA/QC tolerances:

- The completeness of the Laboratory Reports, verifying that all required components of the report are present and that the samples indicated on the accompanying chain-ofcustody are addressed in the report.
- The calibration and tuning records for the laboratory instruments used for the sample analyses.
- The results of internal standards analyses.
- The results of laboratory blank analyses.
- The results of LCS analyses.
- The results of MS/MSD analyses.
- Compound identification and quantification accuracy.
- Laboratory precision, by reviewing the results for blind field duplicates.

Weston Solutions, Inc. - Quality Assurance Sampling Plan, Jackpile-Paguate Uranium Mine, Paguate, Cibola County, NM

6. QUALITY ASSURANCE

Quality assurance will be conducted in accordance with the WESTON Corporate Quality Management Manual, dated March 2004; the WESTON START-3 Quality Management Plan, dated August 2007; and EPA Quality Assurance/Quality Control Guidance for Removal Activities, dated April 1990. Following receipt of the TDD from EPA, a Quality Control (QC) officer is assigned and monitors work conducted throughout the entire project including reviewing interim report deliverables and field audits. The START-3 PTL will be responsible for QA/QC of the field investigation activities. The designated laboratory utilized during the investigation will be responsible for QA/QC related to the analytical work. START-3 will also collect samples to verify that laboratory QA/QC is consistent with the required standards and to validate the laboratory data received.

6.1 SAMPLE CUSTODY PROCEDURES

Because of the evidentiary nature of sample collection, the possession of samples must be traceable from the time the samples are collected until they are introduced as evidence in legal proceedings. After sample collection and identification, samples will be maintained under chain-of-custody (COC) procedures. If the sample collected is to be split (laboratory QC), the sample will be allocated into similar sample containers. Sample labels completed with the same information as that on the original sample container will be attached to each of the split samples. All personnel required to package and ship coolers containing potentially hazardous material will be trained accordingly.

START-3 personnel will prepare and complete chain-of-custody forms using the Scribe Environmental Sampling Data Management System (SCRIBE) for all samples sent to a START-3 designated off-site laboratory. The chain-of-custody procedures are documented and will be made available to all personnel involved with the sampling. A typical chain-of-custody record will be completed each time a sample or group of samples is prepared for shipment to the laboratory. The record will repeat the information on each sample label and will serve as documentation of handling during shipment. A copy of this record will remain with the shipped samples at all times, and another copy will be retained by the member of the sampling team who

originally relinquished the samples. At the completion of the project, the data manager will export the SCRIBE chain-of-custody documentation to the Analytical Service Tracking System (ANSETS) database.

Samples relinquished to the participating laboratories will be subject to the following procedures for transfer of custody and shipment:

- Samples will be accompanied by the COC record. When transferring possession of samples, the individuals relinquishing and receiving the samples will sign, date, and note the time of the sample transfer on the record. This custody records document transfer of sample custody from the sampler to another person or to the laboratory.
- Samples will be properly packed for shipment and dispatched to the appropriate laboratory for analysis with separate, signed custody records enclosed in each sample box or cooler. Sample shipping containers will be custody-sealed for shipment to the laboratory. The preferred procedure includes use of a custody seal wrapped across filament tape that is wrapped around the package at least twice. The custody seal will then be folded over and stuck to seal to ensure that the only access to the package is by cutting the filament tape or breaking the seal to unwrap the tape.
- If sent by common carrier, a bill of lading or airbill will be used. Bill of lading and airbill receipts will be retained in the project file as part of the permanent documentation of sample shipping and transfer.

SOPs 1101.01 and 1102.01 describe these procedures in more detail.

6.2 PROJECT DOCUMENTATION

All documents will be completed legibly in ink and by entry into field logbooks and SCRIBE. Response Manager will be used after direction of the EPA SAM.

6.2.1 Field Documentation

The following field documentation will be maintained as described below.

Field Logbook

The field logbook is a descriptive notebook detailing site activities and observations so that an accurate, factual account of field procedures may be reconstructed. All entries will be signed by the individuals making them. Entries should include, at a minimum, the following:

Jackpile-esi oasp.doc 6-2 cerclis id nmn000607033

- Site name and project number.
- Names of personnel on-site.
- Dates and times of all entries.
- Description of all site activities, including site entry and exit times.
- Noteworthy events and discussions.
- Weather conditions.
- Site observations.
- Identification and description of samples and locations.
- Subcontractor information and names of on-site personnel.
- Dates and times of sample collections and chain-of-custody information.
- Records of photographs.
- Site sketches.
- Calibration results.

Sample Labels

Sample labels will be securely affixed to the sample container. The labels will clearly identify the particular sample and include the following information:

- Site name and project number.
- Date and time the sample was collected.
- Sample preservation method.
- Analysis requested.
- Sampling location.

Chain-of-Custody Record

A chain-of-custody will be maintained from the time of sample collection until final deposition. Every transfer of custody will be noted and signed for, and a copy of the record will be kept by each individual who has signed it.

Custody Seal

Custody seals demonstrate that a sample container has not been tampered with or opened. The individual who has custody of the samples will sign and date the seal and affix it to the container in such a manner that it cannot be opened without breaking the seal.

Photographic Documentation

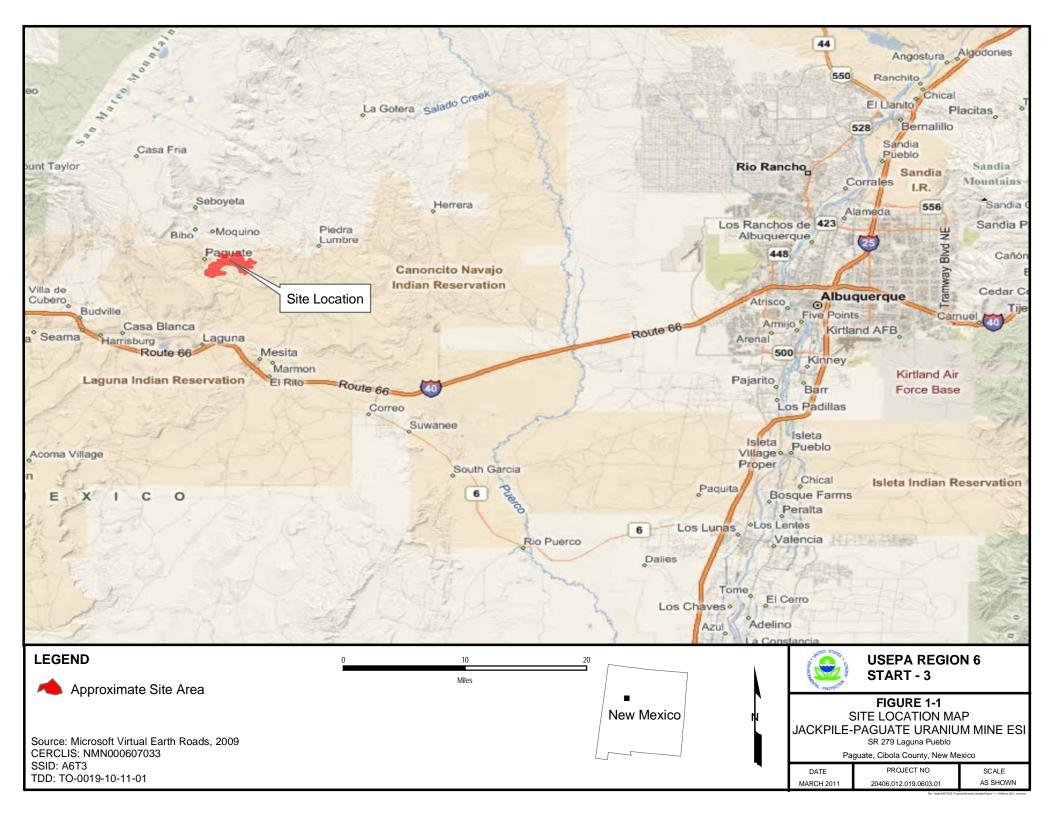
START-3 will take photographs to document site conditions and activities as site work progresses. Initial conditions should be well documented by photographing features that define

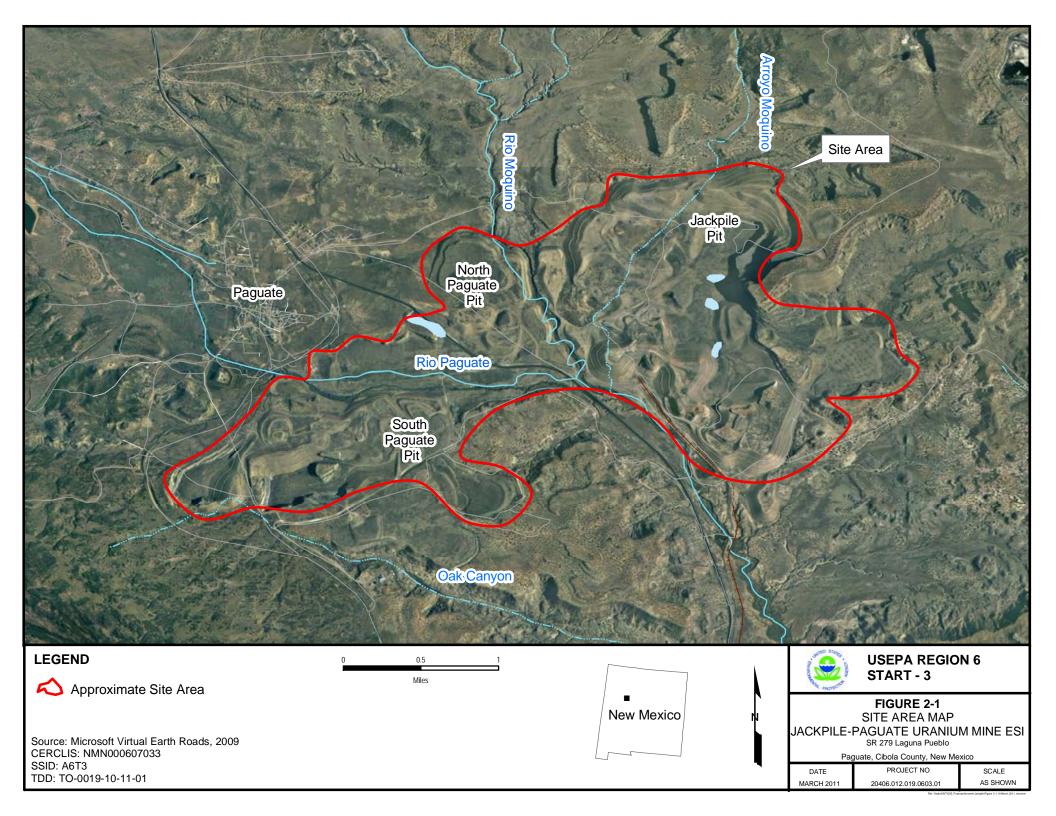
the site-related contamination or special working conditions. Representative photographs should be taken of each type of site activity. The photographs should show typical operations and operating conditions as well as special situations and conditions that may arise during site activities. Site final conditions should also be documented as a record of how the site appeared at completion of the work.

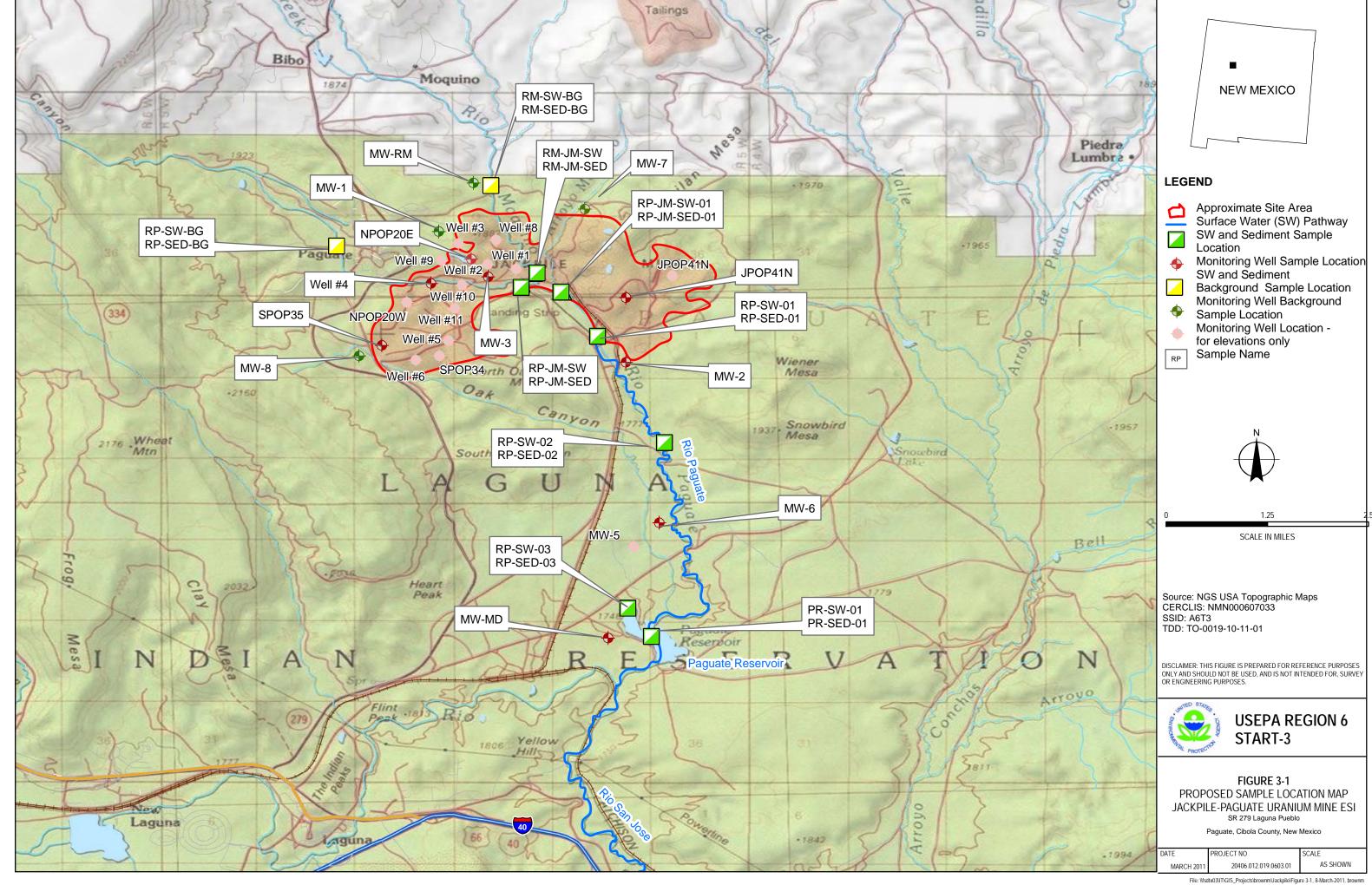
All photographs should be taken with either a film camera or digital camera capable of recording the date on the image. Each photograph will be recorded in the logbook with the location of the photographer, direction the photograph was taken, the subject of the photograph, and its significance (i.e., why the picture was taken).

6.2.2 Report Preparation

At the completion of the project, START-3 will review and validate all laboratory data and prepare a draft report of field activities and analytical results for EPA SAM review. Draft deliverable documents will be uploaded to the EPA TeamLink website for EPA SAM review and comment.







APPENDIX A
EPA CONTRACT LABORATORY PROGRAM GUIDANCE FOR FIELD SAMPLERS



Office of Superfund Remediation and Technology Innovation



Contract Laboratory Program Guidance for Field Samplers

Foreword

The intent of the Contract Laboratory Program (CLP) Guidance for Field Samplers is to replace the CLP Samplers Guide. This guidance document is designed to provide users with general information regarding environmental sample collection for the United States Environmental Protection Agency's (USEPA) Contract Laboratory Program (CLP). This document provides minimum CLP requirements, an explanation of the general sampling process sequence of events, and any related information. The appendices contain useful reference information and checklists to aid in planning and documenting sampling activities.

CLP users also are encouraged to review the Introduction to the Contract Laboratory Program document that contains a general overview of the CLP, how it works, and how to access the program. The CLP requires samplers to use the functionality provided by the Field Operations Records Management System (FORMS) II LiteTM software, which is the preferred means of creating CLP sample documentation. For guidance in using the software to record and submit sampling data, users should reference the FORMS II Lite User's Guide.

Both the Introduction to the Contract Laboratory Program and the Contract Laboratory Program Guidance for Field Samplers can be downloaded from the CLP Web site at the following address:

http://www.epa.gov/superfund/programs/clp/guidance.htm

The FORMS II Lite User's Guide can be downloaded from the CLP Web site at the following address:

http://dyncsdao1.fedcsc.com/itg/forms2lite/doc.html

For more information regarding the CLP or this guide, please contact Elizabeth Holman via email at Holman.Elizabeth@epa.gov or via telephone at (703) 603-8761.

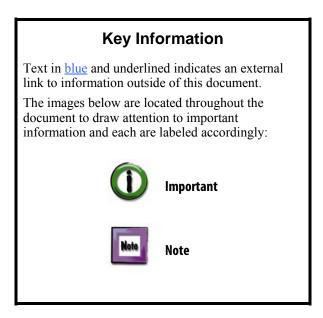


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List of Acronyms

ASB Analytical Services Branch

CERCLA Comprehensive Environmental Response, Compensation, and Liability Act

CLP Contract Laboratory Program

CLP Project Officer

CRQL Contract Required Quantitation Limit
CVAA Cold Vapor Atomic Absorption
DOT Department of Transportation
DQO Data Quality Objective

dbfDatabase FileETEastern Time

FORMS II Lite[™] Field Operations Records Management System II Lite

FSP Field Sampling Plan
HCN Hydrocyanic acid

IATA International Air Transport Association

ICP-AES Inductively Coupled Plasma-Atomic Emission Spectroscopy

ICP-MS Inductively Coupled Plasma-Mass Spectrometry

MS Matrix Spike

MSD Matrix Spike Duplicate
NAHSO₄ Sodium Bisulfate
NPL National Priorities List
OSC On-scene/on-site Coordinator

OSHA Occupational Safety and Health Administration

OSRTI Office of Superfund Remediation and Technology Innovation

OSWER Office of Solid Waste and Emergency Response

PCBs Polychlorinated Biphenyls
PE Performance Evaluation
PM Program Manager
ppb Parts-Per-Billion
ppt Parts-Per-Trillion

PRP Potentially Responsible Party

PT Proficiency Testing
PTFE Polytetrafluoroethylene
PVC Polyvinyl Chloride
QA Quality Assurance

QAPP Quality Assurance Project Plan

QASPER Quality Assurance Sampling Plan for Environmental Response

QATS Quality Assurance Technical Support

QC Quality Control

RAS Routine Analytical Services **RPM** Remedial Project Manager

RSCC Regional Sample Control Center Coordinator

RSM Regional Site Manager
SAM Site Assessment Manager
SAP Sampling Analysis Plan

SARA Superfund Amendments and Reauthorization Act

SDGSample Delivery GroupSMCSystem Monitoring CompoundSMOSample Management OfficeSOPStandard Operating Procedure

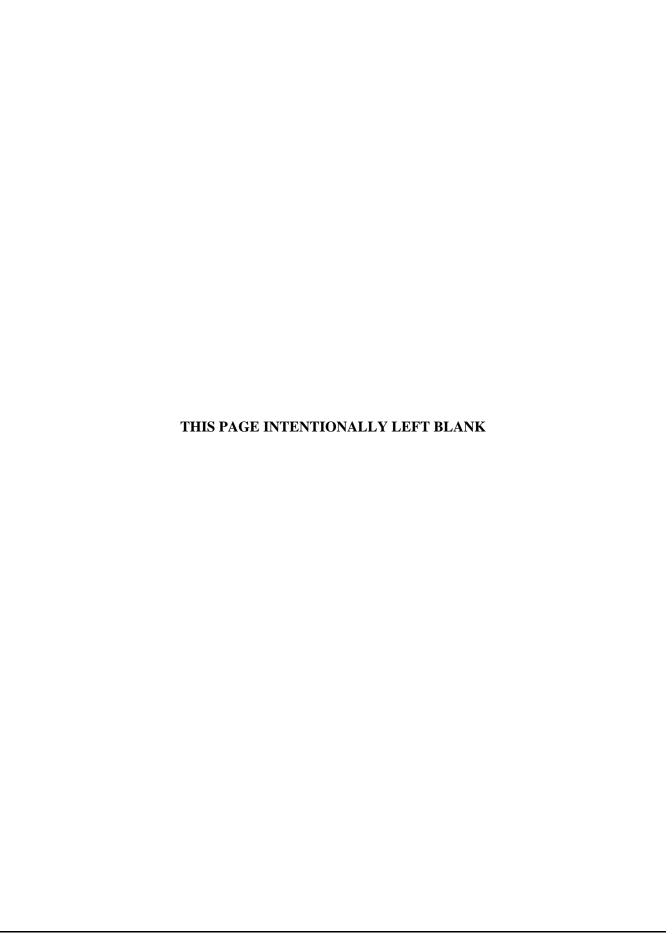
SOW Statement of Work

SVOA Semivolatile Organic Analyte
TR/COC Traffic Report/Chain of Custody

txt Text File United Nations

USEPA United States Environmental Protection Agency

VOA Volatile Organic Analyte
XML eXtensible Markup Language



1.0 INTRODUCTION

1.1 About this Guide

This document describes the important organizational roles and responsibilities for those who plan and conduct environmental sample collection projects for analysis through the Superfund's Contract Laboratory Program (CLP). This chapter introduces the structure and purpose of this document. Chapter 2, *Pre-field Activities*, addresses pre-field planning activities that the sampling team could complete prior to the actual sampling event. Chapter 3, *In-field Activities*, addresses those activities that need to be completed during the sampling event.

Appendix A describes the functions within a sampling project which are taken from the Quality Assurance Project Plan requirements. Appendix B and Appendix C contain the sample collection guidelines for Volatile Organic Analytes (VOAs) in soil and in water. Appendix D recommends sampling techniques. Appendix E contains checklists to help the sampler ensure that all necessary steps are completed.



A project and site-specific Quality Assurance Project Plan (QAPP) providing Regional guidance will override guidance given within this document.

1.2 Overview of the CLP

The CLP is a national program of commercial laboratories under contract to support the USEPA's nationwide effort to clean up designated hazardous waste sites by supporting its Superfund program. The Superfund program was originally established under the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) of 1980 and presently exists under the Superfund Amendments and Reauthorization Act (SARA) of 1986.

The CLP uses state-of-the-art technology to provide users with analytical services. The program provides data of known and documented quality to support USEPA enforcement activities or other user needs. To achieve this goal, the CLP has established strict Quality Control (QC) procedures and detailed documentation requirements. Current CLP users include the USEPA Regions, States and Tribal governments, and other Federal agencies. CLP users also are encouraged to review the *Introduction to the Contract Laboratory Program* document that contains a general overview of the CLP, how it works, and how to access the program.

1.2.1 Key Players Within the CLP

In coordinating Superfund sampling efforts, the Analytical Services Branch (ASB) is supported by the Sample Management Office (SMO) contractor, the Regional CLP Project Officers (CLP POs), the Regional Sample Control Center Coordinators (RSCCs), and the Regional Site Managers (RSMs), including Site Assessment Managers (SAMs), On-scene/On-site Coordinators (OSCs), and Remedial Project Managers (RPMs). Samplers may work directly with the RSCC and/or RSM (or equivalent), and/or an OSC from the Field Support Section during a sampling event. See Table 1-1 for a brief description of the functions performed by key participants (functions may vary by Region).

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Table 1-1. Participants in the CLP Sampling Process

	tote 1-1. Participants in the CLP Sampling Process
Participants	Responsibilities
Analytical Services Branch	USEPA ASB directs the CLP from within the Office of Superfund Remediation and Technology Innovation (OSRTI) in the Office of Solid Waste and Emergency Response (OSWER). ASB responsibilities include:
	 Development of the Statements of Work (SOWs) that define required analytical methods (including QC, detection/quantitation limits, and holding times) for the analytical services procured under the CLP;
	 Development and implementation of policies and budgets for Superfund analytical operations;
	 Development of information management policies and products for analytical data;
	 Management of SMO and Quality Assurance Technical Support (QATS) contracts;
	 National administration, evaluation, and management of the CLP; and
	 Direction of CLP Quality Assurance (QA) activities in coordination with overall OSWER QA activities.
	To obtain the most current ASB contact list, refer to the following Web site:
	http://www.epa.gov/superfund/programs/clp/contacts.htm#ASB
CLP Sample Management Office	The contractor-operated SMO provides necessary management, operations, and administrative support to the CLP. SMO receives Regional analytical requests, coordinates and schedules sample analyses, and tracks sample shipments. SMO also receives and checks data for completeness and compliance, processes laboratory invoices, and maintains a repository of sampling records and program data.
CLP Contract Laboratories	The contractor-operated laboratories within CLP provide necessary analytical services for the isolation, detection, and quantitation of the CLP's target compounds and analytes.
Regional CLP Project Officer	The CLP PO monitors the technical performance of the contract laboratories in each Region. The CLP PO works closely with ASB Program Managers (PMs) to identify and resolve laboratory technical issues, and leads laboratory on-site evaluations. To obtain the most current CLP PO contact list, refer to the following Web site:
	http://www.epa.gov/superfund/programs/clp/polist.htm
Regional Sample Control Center Coordinator	In most Regions, the RSCC coordinates sampling efforts and serves as the central point-of-contact for sampling questions and problems. The RSCC works with SMO to schedule sample shipments to laboratories. In addition, the RSCC's activities may include: informing SMO of sample shipment, cancellations, special instructions, and sampling issues. To obtain the most current RSCC contact list, refer to the following Web site: http://www.epa.gov/superfund/programs/clp/rscclist.htm
Regional Site Manager	The RSM Coordinates the development of acceptance or performance criteria and oversees project-specific contractors, state officials, or private parties conducting site sampling efforts. The RSM could be the SAM, the OSC, or the Remedial Project Manager (RPM).
Field Support Section	The Field Support Section consists of personnel such as the OSC, SAM, and RPM. In most Regions, the Field Support Section develops Standard Operating Procedures (SOPs) for field sampling and related procedures, and assists sampling teams in following those SOPs. The sampling team determines what type(s) of CLP services will be required for a particular sampling event. The Field Support Section reviews Sampling Analysis Plans (SAPs) prepared by sampling teams and oversees sampling teams in the field. The Field Support Section may also prepare their own SAPs, perform sampling activities in the field, and analyze and report the results of their sampling events to the RSM.

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1.3 Overview of the Sampling Process

Once USEPA has determined that physical, chemical, and/or biological testing of a site is necessary, samples of material from the site area must be collected. The type of material that must be collected and the analytical method to be used depends upon the physical location of the site, detection level(s), site history (previous sampling), and known or unknown conditions and contaminants. The sampling process includes carefully planned and consistently applied procedures that produce accurate and legally defensible data. The sampling team should consider the procedures and plans presented in this guide as minimum sampling process guidelines to

At-a-Glance: Overview of the Sampling Process

- ✓ Procedures must be consistent.
- ✓ Analytical data must be accurate and defensible.
- ✓ Procedures must meet minimum requirements.

maintain sample integrity and identity. Samples should be collected according to the approved project and site-specific QAPP and SAP. This document does not define specific sampling procedures because specific sampling protocols depend on individual site conditions, Regional requirements, and acceptance and performance criteria. Since Regions may have their own specific requirements for individual sampling programs, they are responsible for generating Region-specific sampling SOPs.

1.3.1 Procedures Must be Consistent

The purpose of sampling is to collect representative portions from a suspected contaminated site. Sample collection is critical to determining the presence, type, concentration, and extent of environmental contamination by hazardous substances, thus it is a crucial part of every sampling and environmental testing effort. Sampling procedures must be consistently written and followed to mitigate risk of error and the expense of re-sampling.

Failure to follow proper sampling and shipping procedures could result in samples that are contaminated, broken, mislabeled, lost during shipping, or unusable because of a missed holding time. If procedures are inconsistently or improperly followed, any resultant analytical data may be inaccurate and may not be defensible in a court of law.



If re-sampling is needed due to improper sampling, the sampling team may incur the cost.

1.3.2 Analytical Data Must be Accurate and Defensible

The data gathered during sampling activities helps to accurately characterize contaminated waste sites so that the impact on human health and the environment can be properly evaluated. Acquiring accurate and defensible data that will be accepted in a court of law is the CLP's primary objective; therefore, the sampler must collect samples according to strict sampling procedures, plans, and guidelines. USEPA and many other Federal agencies use data resulting from analytical testing of soil/sediment/aqueous samples to:

- Determine if a site is contaminated with organic and/or inorganic compounds;
- Identify pollution sources and Potentially Responsible Parties (PRPs);
- Validate remedial design methodologies;
- Assess response and remedial priorities;
- Assess risk to human health and the environment;
- Determine appropriate cleanup actions; and
- Determine cleanup achievements.

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1.3.3 Sampling Procedures and Guidelines Must Meet Minimum Requirements

It is imperative that samplers be aware of the minimum CLP and Regional requirements that directly impact and define how a sampling event will take place. It is important to note that the procedures and guidelines set forth in this document are considered minimum CLP requirements. Samplers should reference the following sections within this document that specifically address important requirements that must be met for a successful sampling event:

- Section 1.4.1 CLP Documentation Requirements;
- Section 2.4.1 Request Scheduling of Analysis, SMO-assigned Case Numbers, CLP Sample Numbers, and Laboratory Contact Information;
- Section 2.7 Comply with Transportation and Shipping Requirements;
- Section 2.8 Provide Shipment Notification;
- Section 3.1 Collect Samples; and
- Section 3.2 Complete Documentation.

1.4 Overview of Sampling Documentation Requirements

The sampler must properly document samples collected for analysis in order to uniquely identify each sample and ensure adequate chain-of-custody procedures. When collecting samples, the sampler should always keep in mind that any samples collected may be used in future litigation. This is especially important when samples are from privately owned property. If sampling on privately owned property, samplers should also provide the property owner with a receipt for samples collected and removed from that owner's property. Samplers may also be required by a Region to use a sample label, sample tag, or field operations records documenting information such as daily activities, equipment and materials used, personnel involved, site security, etc. These types of documentation help ensure proper sample identification and provide additional chain-of-custody records.

The documentation required by a Region for a sampling event is outlined in project plans such as the QAPP, SAP, and Field Sampling Plan (FSP).

At-a-Glance: Overview of the Sampling Document Requirements

- Must use FORMS II Lite to create sample documentation. Analytical data must be accurate and defensible.
- ✓ CLP documentation requirements:
 - CLP Sample Number
 - SMO-assigned Case Number
 - Traffic Report/Chain of Custody (TR/COC) Record
 - Sample Labels
 - Sample Tags
 - Custody Seals
 - Field Operation Records



Under no circumstances should the site name appear on any documentation that is sent to the laboratory (for the CLP).

1.4.1 CLP Documentation Requirements

Samplers must:

- 1) Record the CLP Sample Number on each sample bottle;
- 2) Complete the Traffic Report/Chain of Custody (TR/COC) Record using the FORMS II Lite software, making sure to indicate on the TR/COC Record if the samples require the use of a Modified Analysis;
- 3) Complete and attach sample labels;
- 4) Complete and attach sample tags to meet Regional requirements;
- 5) Complete and attach custody seals to meet Regional requirements; and
- 6) Complete field operations records, as necessary.

Please contact your RSCC (see Table 1-1) for information regarding CLP Sample Numbers, SMO-assigned Case Numbers, TR/COC Records, and chain-of-custody seals for sampling events.

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For information regarding using FORMS II Lite to create and complete a TR/COC Record, refer to the following Web site:

http://www.epa.gov/superfund/programs/clp/f2lite.htm

1.4.1.1 CLP Sample Number

A CLP Sample Number is unique per sampling location and is used to identify and track samples throughout the sampling and analytical processes and is recorded on many types of sampling documentation (e.g., TR/COC Records, sample labels, and sample tags). CLP Sample Numbers are provided to samplers by their RSCC or SMO.

Samplers must contact their RSCC (or their designee) to obtain CLP Sample Numbers for their sampling event. Samplers must correctly assign the CLP Sample Numbers to the appropriate sample bottle or container. Please refer to Section 3.2.1 for more detailed information regarding the use of CLP Sample Numbers.



If the sampler has any questions regarding the assignment of CLP Sample Numbers, they should contact their RSCC.

1.4.1.2 SMO-assigned Case Number

SMO-assigned Case Numbers are used to track groups of samples throughout the sampling and analytical processes and are recorded on many types of sampling documentation (e.g., TR/COC Records, sample labels, and sample tags). Samplers must correctly assign the SMO-assigned Case Number to the appropriate sample bottle or container. To obtain a SMO-assigned Case Number, samplers must contact their RSCC (or their designee).

1.4.1.3 Laboratory Assignment

Samplers are responsible for shipping samples to the appropriate SMO-assigned laboratory for analysis. Samplers must contact their RSCC (or their designee) to obtain their laboratory assignment or they may be provided by SMO.

1.4.1.4 TR/COC Record

The TR/COC Record is used as physical evidence of sample custody and functions as a permanent record of each sample collected.

Per CLP documentation requirements, each cooler must contain a TR/COC Record that lists all the samples contained therein.

In an effort to automate sample documentation in the field, ASB has developed a stand-alone, Windows-based software application that samplers can use to automatically create and generate sample documentation. The FORMS II Lite software allows users to enter information prior to and during sampling events. It allows users to multi-task and electronically create, edit, and print documentation associated with sampling activities. Users can customize data entry screens throughout the entire documentation process. Users can also customize the format and content of sample labels based on specific requirements.

The program simplifies and accelerates the tedious manual sample documentation process by reducing the generation of handwritten documents by almost 70%. The FORMS II Lite software enables samplers to:

- Increment CLP Sample Numbers or manually assign their own unique, project-specific non-CLP Sample Numbers;
- Input the SMO-assigned Case Number into the appropriate field;
- Create sample labels, sample tags, TR/COC Records, Sample Weight forms, and receipts for samples taken from a site;
- Track samples from the field to the laboratory;

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- Electronically capture sample information into databases; and
- Export electronic data as a database File (.dbf), Text (.txt), or eXtensible Markup Language (.xml) file.

USEPA requires samplers to use the FORMS II Lite software for all CLP sampling efforts. For assistance with obtaining or using the FORMS II Lite software, please contact the FORMS II Lite Help Desk at 703-818-4200 from 9:00 AM - 5:00 PM Eastern Time (ET). For additional information regarding FORMS II Lite use and training, please refer to the following Web site:

http://www.epa.gov/superfund/programs/clp/f2lite.htm

1.4.1.5 Chain-of-Custody Seals

A chain-of-custody seal is any adhesive label or tape that can be used to seal a sample bottle, container, plastic bag, or shipping cooler such that if it is opened or tampered with, the seal will be broken. Custody seals must be placed on each sample bottle, container, or bag (as appropriate) and each shipping cooler or container. The custody seal is an excellent means of maintaining a record of chain-of-custody, as well as guarding against possible sample contamination or tampering during shipping.

1.4.1.6 Sample Labels

A sample label is a sticker attached to a sample bottle or container that contains a sample. Sample labels are affixed to each sample container as samples are collected in the field or affixed prior to going in the field. A sample label must contain, at a minimum, a CLP Sample Number so that they can be associated with, and listed on, the associated TR/COC Record. The sample label may also include the required analysis/fraction and preservative used (to eliminate confusion at the laboratory). Samplers should refer to their project plans for Region-specific sample label requirements.

1.4.1.7 Sample Tags

A sample tag identifies a sample bottle or container that contains a sample. The tag also provides specific analytical direction and proof that a sample existed. To support the use of sample data in potential enforcement actions, samples with other than in situ measurements (e.g., pH, temperature, conductivity) can be identified with a sample tag. A CLP Sample Number and SMO-assigned Case Number must be recorded on a sample tag to indicate that the sample container comprises the whole sample in the case where there is just one container of sample, or part of the indicated sample in the case of multiple containers of sample. Samplers should refer to their project plans for Region-specific sample tag requirements.

1.4.1.8 Field Operation Records

Samplers should maintain complete, accurate, and legible field operations records as they perform a sampling activity. The following records are included: Field Logbooks; Corrective Action Reports; Sampling Trip Reports; supplemental standardized forms; logs; and records such as maps or photographs that document each step of the work performed in the field. Samplers should refer to their project plans for Region-specific field operations record requirements. These records are very important tools because they are considered part of the official project file when legal issues arise.

1.4.1.9 Weight Logs

A sample weight log identifies the tared, sample and final weights per bottle for VOA samples. In order to support Method 5035 for VOAs, samplers should enter tared and final weights per bottle in the CLP Sample Weight Log.

2.0 PRE-FIELD ACTIVITIES

This chapter provides instructions for completing the suggested prefield activities that samplers could complete prior to performing sampling activities. These important pre-field activities will save time and help the sampler to better prepare for the sampling event. Samplers should be aware of issues routinely arise during the sampling process so that samplers can avoid making the same mistakes or having the same problems that could adversely affect their sampling event. Samplers are also expected to review all pertinent project plans and meet both CLP and Regional requirements that directly impact the structure and purpose of a sampling event.

The project plans provide information such as the types and numbers of samples to be collected, the analytical methods to be used based on the desired level of quantitation, and the necessary equipment and supplies. The plans also describe the sampling method which may require different specific sample volumes/masses, containers,

preservation, shipping, and handling to maintain the integrity of the samples without degradation or contamination.

At-a-Glance: Pre-field Activities

- Prepare for and communicate during a sampling event.
- Review project plans containing Regional requirements.
- ✓ Plan to meet documentation requirements.
- ✓ Obtain any necessary permits, licenses, and clearances.
- ✓ Identify and obtain sampling materials.
- ✓ Comply with transportation and shipping requirements.
- ✓ Provide shipment notification.
- ✓ Perform Readiness Review/Run-through.

In addition to reviewing project plans, samplers should determine if the sampling site is privately or publicly owned and obtain the necessary permission to access the sampling site. If the site is privately owned, samplers should make sure to have receipts for available samples to provide to the owner for all samples collected and removed from their property. Samplers must also prepare to identify and obtain sampling materials, prepare to meet documentation requirements by obtaining and learning to use the required software, comply with transportation and shipping requirements, and perform a readiness review/dry run of the sampling process.

2.1 Prepare for a Sampling Event

Samplers must prepare to meet CLP and Regional requirements for a sampling event, appropriately use the CLP Sample Number and SMO-assigned Case Number, complete the TR/COC Record using the FORMS II Lite software, and complete and attach the custody seal(s). It is very important that the sampler include the correct CLP Sample Number on each sample. It is also imperative that the TR/COC Record be accurately completed and submitted with the sample(s). Finally, the sampler must accurately and legibly complete and attach a custody seal to each sample container, or plastic sample bag (as appropriate), and each shipping cooler or container.

However, meeting the sampling requirements requires more than just the proper application of a CLP Sample Number on each sample, completion of the TR/COC Record, and use of a custody seal. The actual collection of samples, packaging, and shipping of those samples are equally important to a successful sampling event.

For example, if a sampler collects an insufficient volume of a sample, the laboratory may not be able to perform the requested analysis. Insufficient sample volumes may also result in a laboratory being unable to perform laboratory quality control, such as Matrix Spike (MS), Matrix Spike Duplicate (MSD), and Duplicate sample analysis. Additionally, if the laboratory receives a sample that is either unpreserved or the sample pH is outside of the required range, the sample cannot be properly analyzed.

Unfortunately, improper shipping and labeling processes and procedures often result in:

- Samples being shipped to the wrong laboratory;
- Broken or empty samples being received at the laboratory; and
- Custody seals or sealant tape that is missing or broken on sample bottles, containers, plastic bags, or shipping coolers shipped to the laboratories.

The importance of completing the paperwork associated with a sampling event cannot be overemphasized. Samplers must make a conscientious effort to accurately complete the TR/COC Record since this is the main document used to derive vital information about a particular sample. The person completing a TR/COC Record

must be careful to avoid errors such as the appropriate sample(s) not being listed, or the wrong samples being listed. In an effort to eliminate such errors and the confusion that can be associated with handwritten TR/COC Records, samplers must use the FORMS II Lite software to complete the TR/COC Record and other associated sampling documentation.

It is extremely important that QC samples, including field sample duplicates, field samples for Matrix Spike and Matrix Spike Duplicate analyses, and Proficiency Testing (PT) samples, also known as Performance Evaluation (PE) samples, be designated and labeled per Regional guidance by samplers in the field. Mislabeling of QC samples can result in improper and/or inaccurate analysis of a sample at the laboratory.

2.2 Communicate During a Sampling Event

Communication is a key element in planning, administrating, and conducting a sampling event. It is extremely important that all parties involved in a sampling event be in contact throughout the sampling process. The procedures and recommendations outlined in this guide are based on more than 20 years of experience. It has been demonstrated that approximately 50% of all sampling efforts have been negatively affected by incorrect sampling procedures and poor communication among participants.

The key elements of communication for a sampling event include the relationship between the RSCC, SMO, the samplers in the field, and the laboratories who will be accepting the samples. For instance, the samplers must contact the RSCC to start the process for setting up a sampling event. The RSCC will in turn contact SMO who will schedule the sampling event, establish laboratory availability, and arrange for the laboratory to accept projected samples. SMO will then communicate the laboratory assignment to the Region and possibly the sampler.



The sampler should contact the RSCC (per Regional guidelines) and allow enough time for the RSCC to contact SMO at least a week prior to the sampling event.

SMO provides SMO-assigned Case and CLP Sample Numbers in time for the sampling event. SMO also schedules a laboratory and makes sure the laboratory will not have any capacity problems. Communication is also important because if there is a change in the sampling event due to a cancellation or an increase or decrease in the number of samples that will be sent to the laboratory, the sampler can contact the RSCC who can work with SMO to remedy potential capacity, availability, or overbooking problems.

2.3 Review Project Plans Containing Regional Requirements

In addition to meeting CLP requirements, the sample collection process must fulfill numerous Regional requirements. These requirements are determined by a variety of factors that affect how samples should be collected for an individual sampling event. These factors include:

- The type of samples being collected (organic/inorganic, water, soil/sediment, etc.);
- The method by which the samples will be analyzed;
- The acceptance or performance criteria (i.e., Data Quality Objectives [DQOs]); and
- The type of data needed.

The QAPP for each sampling project is written to meet requirements outlined in the documents *EPA Requirements* for Quality Assurance Project Plans (QA/R-5), EPA Guidance on Quality Assurance Project Plans (G-5), and Regional QAPP preparation documents. The QAPP is prepared in advance of field activities and is used by samplers to develop any subsequent plans such as the Sampling SAP or the FSP. Samplers should review the QAPP and any subsequent project plans for information outlining the basic components of a sampling activity. QAPP and project plans should be finalized and approved by appropriate Regional QA personnel, the OSC, SAM, or the RPM before providing them to the sampling team. This should be done prior to the start of field activities. Appendix A explains the functions within a sampling project (as these functions relate to a sampling event) and the elements of that function as described in a typical QAPP. Copies of all project plans and relevant SOPs should be maintained in the field for the duration of the sampling project.

2.4 Plan to Meet Documentation Requirements

Sampling events require a variety of accurate and complete documentation. Samplers should review their project plans to determine the types of documentation that must be completed for a sampling project and to ensure that the appropriate documentation will be on-hand in the field. The CLP documentation requirements include the CLP Sample Number, the SMO-assigned Case Number, the TR/COC Record, sample labels, sample tags, custody seals, and field operations records (as necessary). Samplers need to request SMO-assigned Case and CLP Sample Numbers for each

At-a-Glance:

Plan to meet documentation requirements.

- Request SMO-assigned Case and CLP Sample Numbers.
- ✓ Prepare sample cooler return documentation.
- ✓ Prepare to use the FORMS II Lite software.

sampling event prior to starting field activities. Samplers also need to make sure that the correct TR/COC Records (Organic TR/COC Record for organic analysis or Inorganic TR/COC Record for inorganic analysis) are being used within the FORMS II Lite software. Finally, samplers should be prepared to complete the appropriate shipping cooler return documentation.

Since samplers are required to use the FORMS II Lite software to prepare and submit sampling project documentation and maintain sample chain-of-custody, software users must be familiar with all emergency back up procedures that should be followed in the event of a system failure. Samplers must have access to FORMS II Litegenerated TR/COC Records at sampling events. If problems are experienced while using the FORMS II Lite software, please contact the FORMS II Lite Help Desk at 703-818-4200 from 9:00 AM - 5:00 PM ET.

In the event of a system crash, samplers must have backup hardcopies of FORMS II Lite TR/COC Records. For information regarding emergency backup procedures, please refer to the following Web site:

http://www.epa.gov/superfund/programs/clp/trcoc.htm

2.4.1 Request Scheduling of Analysis, SMO-assigned Case Numbers, CLP Sample Numbers, and Laboratory Contact Information

SMO-assigned Case Numbers are assigned based on a request for CLP Routine Analytical Services (RAS), which is processed though the RSCC (or his/her designee). The sampler must request the RSCC to schedule CLP RAS analysis. The CLP does have the capacity to schedule sampling on an emergency basis, however the sampler must contact the RSCC (or his/her designee) to obtain details regarding how to handle such a situation. When scheduling a sampling event that will last for more than one week, it is recommended that the sampler contact the RSCC (or his/her designee) on a weekly basis to provide updates. This contact between the sampler, the RSCC (or his/her designee), and SMO is very important because it will ensure better availability of laboratory capacity.

In addition to SMO-assigned Case and CLP Sample Numbers, samplers should make sure to have accurate laboratory contact information, such as:

- Laboratory name;
- Laboratory address;
- Contact name; and
- Laboratory phone number.

This information is used for both TR/COC Records and chain-of-custody documentation and shipping paperwork such as address labels and airbills.

The SMO-assigned Case Number is used to track groups of samples throughout the sampling and analytical processes. Samplers must correctly indicate the assigned Case Number on the appropriate sample bottle or container.



The RSCC (or his/her designee) provides the CLP Case Numbers and Sample Numbers for each sampling event to samplers. Once the CLP Sample Numbers have been provided to the sampler, the sampler can use FORMS II Lite to print them onto sample labels.

The following characters are not to be used in generating CLP Sample Numbers and should never appear on any paperwork submitted to the laboratory: I, O, U, and V.

A CLP Sample Number is defined **as** a number that is unique per sampling location and identifies each CLP sample (see Section 1.4.1.1). Since samples must be identified per analytical program (either organic or inorganic), there are two types of TR/COC Records and two letter codes to denote organic vs. inorganic analysis.

A CLP *sample* is defined as one discrete portion of material to be analyzed that is contained at one concentration level, from one station location for each individual or set of analytical fractions -- provided the fractions are all requested for the same CLP analytical service (i.e., organic or inorganic), and identified by a unique Sample Number.



When samples are collected from several station locations to form a composite sample, the composite sample should be assigned either a number from one of the station locations used during collection, or a unique number that represents the composite sample for tracking purposes. The numbering scheme used internally at a sampling event for identifying composite samples should also be documented appropriately (e.g., in the field logs).

Organic CLP Sample Numbers begin with the Regional letter code, followed by four letters and/or numbers. Inorganic CLP Sample Numbers begin with "M", followed by the Regional letter code and then four letters and/or numbers. See Table 2-1 for Region and letter codes for each sample type (i.e., organic or inorganic).

Dagian	Letter	r Code
Region	Organic	Inorganic
1	A	MA
2	В	MB
3	С	MC
4	D	MD
5	Е	ME
6	F	MF
7	G	MG
8	Н	MH
9	Y	MY
10	J	MJ

Table 2-1. CLP Sample Number Letter Codes

According to CLP guidelines, each individual inorganic water sample may be analyzed for total metals or dissolved metals, but not both. Therefore, water samples collected for total metal and dissolved metal analyses from the same sampling location must be assigned separate (unique) CLP Sample Numbers. A sampler can use the same CLP Sample Number for an inorganic soil or water sample collected for total metals, mercury and cyanide analyses.

Organic soil and water samples may be collected for analysis under the SOM01 SOW to detect:

- Aroclors;
- Semivolatile Organic Analytes (SVOAs);
- Pesticides;
- Volatile Organic Analytes (VOAs); and/or
- Trace Volatile Analytes

Inorganic soil and water samples may be collected for analysis for cyanide, and for metals using Inductively Coupled Plasma-Atomic Emission Spectroscopy (ICP-AES) and Cold Vapor Atomic Absorption (CVAA), under the ILM05.X SOW.

Inorganic water only samples may be collected for analysis for cyanide, and for metals using Inductively Coupled Plasma-Mass Spectrometry (ICP-MS) and CVAA, under the ILM05 SOW.

2.4.2 Prepare Sample Cooler Return Documentation

CLP laboratories must routinely return sample shipping coolers to the appropriate sampling office within 14 calendar days following receipt of shipment from the sampler. For sample coolers to be returned, the

sampler must complete the appropriate cooler documentation and work with Regions and government agencies to provide a cost-effective mechanism for laboratories to return the empty coolers to the appropriate sampling office. The sampling cooler return documentation can be prepared in advance and provided to samplers before field activities begin. The sampler (not the CLP laboratory) is responsible for paying for return of the cooler and should also include shipping airbills bearing the sampler's account number, as well as a return address to allow for cooler return.

To maintain consistency among cooler transportation programs, samplers should:

- Minimize the use of multiple transportation carriers to avoid confusion;
- Use multiple-copy labels so the laboratory and the sampling team can each retain a copy for their records;
- Prepare labels in advance so that the laboratory can simply affix a completed shipping label on the cooler;
- Include third-party billing information (i.e., their shipping account number) on labels so the laboratory will not be billed by the transportation carrier;
- Confirm that the laboratory knows which transportation carrier to use; and
- Include the SMO-assigned Case Number on return information.

2.5 Obtain Municipal Permits, Licenses, and Clearances

Before starting a sampling event, samplers must make sure to obtain the proper municipal permits, accesses to the property, and any government clearances, if required. The sampler must also contact any appropriate utility companies to ascertain where any underground pipes, cables, etc., may be located.

At-a-Glance:

Obtain permits, licenses, and clearances.

- Request access to County, State, Tribal, military, and/or Federal property.
- ✓ Contact private property owner(s).
- ✓ Contact utility companies.

2.5.1 Request Access to County, State, Tribal, Military, and/or Federal Property

Proper access to perform sampling activities is important not only for legal reasons, but also to eliminate delays in work and possible refusal to allow sampling to take place. It is crucial that the appropriate permits, licenses, and clearances be secured to obtain access for sampling activities that will be performed on County, State, Tribal, military, and/or Federal property. The sampler must contact the appropriate government offices or personnel well in advance to determine what kinds of approval are required. Preapproval may be required for specific types of sample collection such as drilling or excavation. For example, drilling on a military base requires pre-approval. Base security may require clearances for all members of the sampling team, including subcontractors. This process may take two or more days.

If arrangements are not made in advance, the team may not be allowed to enter the site until their clearances are processed and the team has been approved to drill. As a result, the sampling schedule is delayed, costing extra time and money.

2.5.2 Contact Private Property Owners

The sampler must obtain written permission from the private property owner(s) before sampling on their property, even if verbal permission has been granted. It is recommended that samplers obtain verbal permission prior to their arrival at the sampling location, but written permission can be obtained on the day of sampling. If a property owner refuses to grant access to their property, it may be necessary for sampling participants to contact the appropriate authorities for assistance.

2.5.3 Contact Utility Companies

The sampler should contact local utility companies (e.g., power, phone, gas, cable, sanitation, etc.) at least one week prior to the sampling event to have underground cables, lines, and pipes flagged and marked. This is required by law. A national one-call directory can be found at:

http://www.digsafelv.com/contacts.htm.

This will eliminate potential safety hazards and service disruption. For example, soil sampling in a residential area may require digging below the soil's surface. It is very important to know where utility lines and pipes are located so that samplers do not hit live electrical wires or rupture gas lines. Samplers should follow Regional or other appropriate program procedures for the procurement of such services. The utility service(s) disruption dates should be confirmed at least two days prior to sampling activities.



Pre-payment of survey fees to local utility companies may be required.

2.6 Identify and Obtain Sampling Materials

Samplers must make sure to be prepared for a sampling project with the appropriate sampling materials (equipment, supplies, sample containers, packing materials, and shipping materials). The equipment and supplies must be properly cleaned, calibrated, and tested as necessary to meet the needs of the sampling project.

At-a-Glance: Identify and obtain sampling materials.

- ✓ Procure appropriate equipment and supplies.
- ✓ Procure sample containers.
- Procure shipping supplies.

2.6.1 Procure Appropriate Equipment and Supplies

Each sampling event requires the procurement of equipment and materials to collect, document, identify, pack, and ship samples. The proper field sampling equipment is vital to a successful sample collection. Regional or other samplers should obtain, and arrange in advance, all of the equipment and supplies required for each sampling event. Samplers should review the project plans to verify that the proper equipment is being used for sample collection.

At a minimum, the following materials are generally required during a sampling event:

- Sample storage containers;
- Packing material;
- Sample containers;
- Shipping containers;
- Access to the FORMS II Lite software for creating sample labels, stickers, tags, and TR/COC Records;
- Custody seals; and
- Sampling equipment such as bowls, augers, pumps, etc.

Sampling events may also require specific items such as:

- Cooler temperature blanks;
- Trip blanks for VOA analysis;
- Preservation supplies (e.g., ice or acid); and
- Specially prepared sample vials (e.g., for SW-846 Method 5035A).

2.6.2 Procure Sample Containers

The analytical protocol(s) to be used for sample analysis often requires the use of a particular type of sample container. The type of container also may depend on the sample matrix and analysis. It is recommended that samplers use borosilicate glass containers, which are inert to most materials, when sampling for pesticides and/or other organics. Conventional polyethylene is recommended when sampling for metals because of the lower cost and absorption rate of metal ions.

Using the wrong container may result in breakage, gathering of an insufficient volume needed to perform sample analysis, or the container material may interfere with the analysis. Therefore, samplers should identify and use the correct sample containers for each sampling event.

Containers procured for a sampling event are usually pre-cleaned and shipped ready-for-use from the manufacturer to the sampling site. Regardless of the type of container used, samplers must ensure that the containers have been analyzed or certified clean to levels below concern for the project. These containers must meet the USEPA container type specifications listed in Table 2-2.

Table 2-2. Container Type Specifications

Reference		Specifications			
Number	Container Type	Closure	Septum		
1	40 mL amber glass vial, 24 mm neck finish.	Polypropylene or phenolic, open-top screw-cap, 15 cm opening, 24-400 size.	24 mm disc of 0.005 in. Polytetrafluoroethylene (PTFE) bonded to 0.120 in. silicone for total a thickness of 0.125 in.		
2	1 L high density polyethylene, cylinder-round bottle, 28 mm neck finish.	Polyethylene cap, ribbed, 28-410 size; F217 polyethylene liner.	N/A		
3	8 oz short, wide mouth, straight-sided, glass jar, 70 mm neck finish.	Polypropylene or phenolic cap, 70-400 size; 0.015 in. PTFE liner.	N/A		
4	4 oz (120 mL) tall, wide mouth, straight-sided, glass jar, 48 mm neck finish.	Polypropylene or phenolic cap, 48-400 size; 0.015 in. PTFE liner.	N/A		
5	1 L amber round glass bottle, 33 mm pour-out neck finish.	Polypropylene or phenolic cap, 33-430 size; 0.015 in. PTFE liner.	N/A		
6	500 mL high density polyethylene, cylinder-round bottle, 28 mm neck finish.	Polypropylene cap, ribbed, 28-410 size; F217 polyethylene liner.	N/A		
7	Coring tool used as a transport device (e.g., 5 g Sampler).	Has built-in closing mechanism.	N/A		
8	250 mL high density polyethylene, cylinder-round bottle, 28 mm neck finish.		N/A		

The information contained in this table is also cross-referenced in the sample collection parameters discussed in Chapter 3. The container Reference Numbers are used in Tables 3-2 and 3-3 under the Containers column. For example, samples collected for low-level soil VOA analysis using SW-846 Method 5035A may require the sampler to use pre-prepared, tared closed-system purge-and-trap vials with a preservative (refer to Appendix B).



Have extra containers readily available for each sampling event in case of breakage, loss, or contamination.

2.6.3 Procure Shipping Supplies

Samples should be correctly packaged into the appropriate shipping containers to reduce the risk of breakage or leakage, and the shipping containers should be appropriately prepared for shipment. Before heading into the field, samplers should refer to the appropriate project plans to determine the types of samples that will be taken during the sampling project so that samplers will have the proper packaging materials at the site for all pertinent samples container types and sample matrices. Samplers should also make sure to obtain the appropriate shipping paperwork (e.g., shipping forms required by the delivery service).

2.7 Comply with Transportation and Shipping Requirements

Samplers are expected to review the applicable project plans to be aware of all State, Federal, Department of Transportation (DOT), and International Air Transport Association (IATA) regulations governing environmental and hazardous sample packaging. The person who ships the samples is responsible for being in compliance with applicable packaging, labeling, and shipping requirements.



Samplers should request and receive sample permits for outside the continental United States, prior to shipping.

Additional information can be obtained on Hazardous Materials Safety Program regulations from the DOT's Research and Special Programs Administration. Federal transportation regulations can be found in 49-CFR Parts 100-185, are available on the Internet at:

http://www.myregs.com/dotrspa/

2.8 Provide Shipment Notification

Some Regions may require that samplers notify their RSCC (or his/her designee) when samples are shipped. Some Regions allow samplers to contact SMO directly to provide shipment notification. It is recommended that samplers contact the RSCC of sample origin to verify if such notification is necessary. If samplers are shipping samples after 5:00 PM ET, samplers must notify the RSCC (or designee) or SMO by 8:00 AM ET on the following business day.



For Saturday delivery at the laboratory, samplers MUST contact the RSCC (or designee) or SMO so that SMO will receive the delivery information by 3:00 PM ET on the Friday prior to delivery.

2.9 Perform Readiness Review/Dry Run

A readiness review/dry run is a test run of the proposed sampling event. This is a recommended practice since it gives samplers a chance to check all plans, documentation software (i.e., FORMS II Lite), and equipment lists for accuracy and completeness prior to sampling activities. It also provides an opportunity to consult with sampling team members to make sure all the elements are in place and everyone understands their tasking before actually going out to the field. Sampling project managers should provide the test or dry run dates and schedules to samplers so that samplers can prepare accordingly.

3.0 IN-FIELD ACTIVITIES

This chapter addresses the in-field activities a sampler will focus on during a sampling event such as: determining the type of samples to be collected; collecting the samples; meeting volume, preservation, and holding time requirements; completing documentation; and packing and shipping samples.

When performing a sampling event, the sampler is expected to follow prescribed sampling techniques. The sampler should also be aware of any special sampling considerations, contamination issues, and sample compositing and mixing methods that could affect their sampling efforts. Please refer to Appendix D for more detailed information.

At-a-Glance: In-field Activities

- ✓ Collecting samples
- ✓ Completing documentation
- ✓ Sampling considerations
- ✓ Procuring shipping supplies



Appropriate Regional guidance and procedures should be consulted for detailed sample collection, preservation, handling and storing, equipment decontamination, and QA/QC procedures.

3.1 Collect Samples

CLP RAS are generally used to analyze samples from Superfund sites. The matrices can be water, soil, or sediment. In some instances, a mixed-matrix sample may be collected which contains either a supernate (for a sediment/soil sample) or a precipitate (for a water sample). In this event, samplers should consult their management plans and/or discuss the required procedures with the RSM or their designee.

A CLP sample consists of all sample aliquots (portions):

- for each individual or set of analytical fractions;
- from one station location;
- for one sample matrix;
- at one concentration level;
- for one laboratory; and
- for one analytical program;

provided that the fractions are all requested from the same CLP analytical service.

In general, it is recommended that two individual samples be collected by separating the aqueous layer from the solid/precipitate layer at the point of collection. They may be assigned two different sample IDs (e.g., Sample IDs ABC124 and ABC125 for Sample ID ABC123), along with a note in the field sample log or tracking system that the sample IDs are derived or related to the same sample ID, to ensure correct follow-up upon receipt of results from the laboratory. Alternatively, they may be assigned the same sample ID, along with a notation of each individual sub-sample or fraction (e.g., Sample IDs ABC123-1 and ABC123-2 or Sample ID ABC123 Fraction 1 and Sample ID ABC123 Fraction 2 for Sample ID ABC123).

3.1.1 Determine Types of Samples to be Collected

Samplers may be required to take several types of samples or sample aliquots during a sampling event. They should refer to their project plans to determine the types of samples or aliquots to be taken, the volumes needed of each sample or aliquot, and the preservation needed for each sample. For an explanation of the various sample types and the requirements for collecting and submitting each particular type, refer to Table 3-1.

Table 3-1. QC Sample Types and CLP Submission Requirements

Sample Type	Purpose	Collection ¹	CLP Sample Number
Field Duplicate	To check reproducibility of laboratory and field procedures. To indicate non-homogeneity.	Collect from areas that are known or suspected to be contaminated. Collect one sample per week or 10% (Regions may vary) of all field samples per matrix, whichever is greater.	Assign two separate (unique) CLP Sample Numbers (i.e., one number to the field sample and one to the duplicate). Submit blind to the laboratory.
Field Blanks	To check cross-contamination during sample collection, preservation, and shipment, as well as in the laboratory. Also to check sample containers and preservatives.	Collect for each group of samples of similar matrix per day of sampling. Organics - Use water (demonstrated to be free of the contaminants of concern). Inorganics - Use metal-free (deionized or distilled) water.	Assign separate CLP Sample Numbers to the field blanks.
Trip Blank (Volatile Organic Analysis Only)	To check contamination of VOA samples during handling, storage, and shipment from field to laboratory.	Prior to going into the field, prepare and seal one sample per shipment per matrix using water demonstrated to be free of the contaminants of concern (deionized water is appropriate). Place this sample in the cooler used to ship VOA samples.	Assign separate CLP Sample Numbers to the trip blanks.
Equipment Blank or Rinsate Blank	To check field decontamination procedures.	Collect when sampling equipment is decontaminated and reused in the field or when a sample collection vessel (bailer or beaker) will be used. Use blank water (water demonstrated to be organic-free, deionized or distilled for inorganics) to rinse water into the sample containers.	Assign separate CLP Sample Numbers to the equipment blanks.
Matrix Spike (MS) and Duplicate (MSD) ² (Organic Analysis Only)	To check accuracy and precision of organic analyses in specific sample matrices.	Collect from areas that are known or suspected to be contaminated. For smaller sampling events (i.e., 20 samples or less), MS/MSD additional volume should be collected in the first round of sampling and included in the first shipment of samples to the laboratory. Collect double or triple volume³ for aqueous samples and soil VOA samples designated for MS/MSD analyses. Additional sample volume is not required for soil samples requiring SVOA, Pesticide, and/or Aroclor analysis. See Appendix B for VOA collection volumes.	Assign the same CLP Sample Number to the field sample and the extra volume for MS/MSD. Identify the sample designated for MS/MSD on the TR/COC Record.
Matrix Spike (MS) and Duplicate (MSD) (Inorganic Analysis Only)	To check accuracy and precision of inorganic analyses in specific sample matrices.	Collect from areas that are known or suspected to be contaminated. For smaller sampling events (i.e., 20 samples or less), Matrix Spike and Duplicates should be collected in the first round of sampling and included in the first shipment of samples to the laboratory. Additional sample volume may be required for inorganic analysis. ⁴	Assign the same CLP Sample Number to the field sample and extra volume (if collected). Identify the sample(s) designated for Matrix Spike and Duplicates on the TR/COC Record.
PE Samples	Specially-prepared QC samples used to evaluate a laboratory's analytical proficiency.	The PE samples contain analytes with concentrations unknown to the laboratory. Designated Regional or authorized personnel (depending on Regional policy) arrange for Case-specific CLP PE samples to be prepared and shipped by the QATS contractor. The PE samples can be shipped to the site, or shipped per Regional direction. QATS provides the appropriate preparation instructions and chain-of-custody materials.	Samplers have no direct interaction with the PE sampling process, but should be aware that such samples do exist within the CLP sampling process. Samplers must, however, order PE samples and ship them to the laboratory if required by the Region.

¹ Consult Regional or Project Manager Guidance for field QC sample frequencies; laboratory QC sample frequencies are generally fixed in the laboratory subcontracts or specified in analytical methods. Current frequency for MS/MSD (organic) and MS/duplicate (inorganic) for the CLP is one sample per twenty field sample of similar matrix.

² Samples sent under the Organic SOW (SOM01) do not require an MS or MSD for Trace VOA, VOA and BNA fractions, but the Region may opt to send them at their discretion.

³ Example of double volume: An aqueous sample for SVOA analysis would require the field sampler to collect at least 2 L of field sample and at least 1 L each for the MS and MSD samples for a total volume of 4 L. If Pesticide or Aroclor MS/MSD analyses are required for the same sample, an additional 4 L must be collected. Double volume is the MINIMUM allowable volume for samples designated for MS/MSD analysis. Triple volume may be sent for MS/MSD samples to allow for sufficient volume for these analyses in the event sample volume is lost as a result of samples breaking, leaking, or laboratory accidents.

⁴ Double volume may be sent for inorganic aqueous MS and MSD samples to allow for sufficient volume for these analyses in the event sample volume is lost as a result of samples breaking, leaking or laboratory accidents.

3.1.1.1 Collect Field QC Samples

Samplers can collect field QC samples and laboratory QC samples to verify that sample quality is maintained during a sampling project.

Field QC samples are designed to assess variability of the media being sampled and to detect contamination and sampling error in the field. The types of field QC samples that are generally collected include field duplicates and field blanks (such as equipment, trip, or rinse blanks). Generally, field duplicate samples should remain "blind" to the laboratory (i.e., they should have separate CLP Sample Numbers).

3.1.1.2 Collect Laboratory QC Samples

A laboratory QC sample is an additional analysis of a field sample, as required by the laboratory's contract. There are three types of laboratory QC samples:

- MS [for organic and inorganic samples];
- MSD [for organic samples only]; and
- Duplicates [for inorganic samples only].



Samplers should obtain Regional guidance regarding the collection of MS and MSD samples (especially for organics analyses).

Samplers should select one sample per matrix per 20 samples as a "laboratory QC" sample. Designated organic laboratory QC samples should be noted on the Organic TR/COC Record. Designated inorganic laboratory QC samples should be noted on the Inorganic TR/COC Record. The laboratory QC sample must not be designated only in the "Field QC Qualifier" column on either the Organic or Inorganic TR/COC Records. Make sure that the laboratory QC sample is included in TR/COC Record samples to be used for the Laboratory QC field.

The sampler should select a field sample as the laboratory QC sample. If the sampler does not select a field sample as the laboratory QC sample, then it is possible that the laboratory could select the field blank (e.g., an equipment or rinsate blank) sample to meet contractual QC requirements. The use of field blanks for laboratory MS/MSD/Duplicate analysis reduces the usability of the data to assess data quality.



In the event of multiple sample shipments during a sampling event, it is recommended that the sampler submit laboratory QC samples in the first sample shipment.

3.1.2 Meet Volume, Preservation, and Holding Time Requirements

Samplers should refer to their project plans to obtain the specific sample volumes to be collected, the preservation needed for those samples, and the technical holding times under which they must submit samples to the scheduled CLP laboratory. Sample collection parameters (to include sample volumes, preservatives, and technical holding times) for organic collection and analysis are listed in Tables 3-2 and 3-3. Sample collection parameters for inorganic analysis and collection are listed in Table 3-4.

3.1.2.1 Collect Sample Volume

Collecting sufficient sample volume is critical. There must be sufficient physical sample volume for the analysis of all required parameters and completion of all QC determinations. The type of analytical procedure(s) to be performed will often dictate the sample volume to collect. For example, each water sample collected for VOA analysis by CLP SOW SOM01 or ILM05 requires a minimum of three vials, each filled completely to a 40 mL capacity. See Appendix C for information regarding the collection of VOAs in water. It is extremely important that samplers refer to their specific project plans to identify and collect the correct sample volume during each sampling event.

When sampling for VOAs in soils, samplers must use SW-846 Method 5035A guidelines included in Appendix B.

3.1.2.2 Preserve Samples

Degradation of some contaminants may occur naturally (e.g., VOAs). The sampler must chemically preserve some water samples for certain analytes before shipping them to the laboratory. The sampler should preserve and immediately cool all samples to 4°C (±2°C) upon collection and samples should remain cooled until the time of analysis (do not freeze water samples). Preservation techniques vary among Regions so the sampler should obtain Region-specific instructions and review the appropriate project plans and SOPs. See Appendix C for information regarding the collection of VOAs in water.

3.1.2.3 Ship within Holding Times

Samplers should ship samples to scheduled CLP laboratories as soon as possible after collection. Daily shipment of samples to CLP laboratories is preferred whenever possible. If samples cannot be shipped on a daily basis, they must be properly preserved and maintained to meet CLP-specified temperatures, holding times, and custody requirements.

The technical holding times are the maximum time allowed between a sample collection and the completion of the sample extraction and/or analysis. In contrast, contractual holding times are the maximum lengths of time that the CLP laboratory can hold the sample prior to extraction and/or analysis. These contractual holding times are described in the appropriate CLP SOW. Contractual holding times are shorter than the technical holding times to allow for sample packing and shipping.



If samplers are shipping samples after 5:00 PM ET, they must notify the RSCC (or designee) or SMO by 8:00 AM ET on the following business day. When making a Saturday delivery, samplers shall contact the RSCC (or designee) or SMO by 3:00 PM ET on the Friday prior to delivery.

Table 3-2. Sample Collection Requirements for CLP SOW SOM01 (VOAs)

Matrix	Container	Sample Type	Minimu	m Number	of Containers	Needed	Minimum Volume/	Important Notes	Preservative	Technical Holding			
Matrix	Туре		with Water	Dry	% Moisture	TOTA L	Mass	-		Time			
	See Table 2- 2, Reference	Samples Only	-	-	-	3		Containers/vials must be filled to capacity with no	Preserve to a pH of 2 with HCl and cool to 4°C (±2°C)				
	Number 1. San SIM	Samples with SIM	-	-	-	4			4		headspace or air bubbles. Refer to Appendix C for	immediately after collection. DO NOT FREEZE water	
Water		Samples with MS/MSD	-	-	-	6	Fill to capacity	samples requiring QC analyses.	samples.	14 days			
	OPTION 1	Samples with SIM and MS/MSD	-	-	-	8							
	OPTION 1 Closed- system Vials	Samples Only	-	3	1	4		Place samples on side prior to being frozen. Refer to Appendix B for	Frozen (-7°C to -15°C) or iced to 4° (±2°C).	14 days			
	See Table 2- 2, Reference Number 1.	Samples with MS/MSD	-	9	1	10	5g	samples requiring QC analyses.		48 hours			
	OPTION 2 Closed-	Samples Only	2	1	1	4		Containers/vials must be filled to capacity with no headspace or air bubbles.	Frozen (-7°C to -15°C) or iced to 4° (±2°C). DO NOT FREEZE water	14 days			
Soil/ Sediment	system contain rights Water See Table 2- 2, Reference Number 1.	Samples with MS/MSD	6	1	5	12	5g	Place samples on side prior to being frozen. Refer to Appendix B for samples requiring QC analyses.	samples.	48 hours			
	OPTION 3 See Table 2- 2, Reference	Samples Only	-	3	1	4		Refer to Appendix B for samples requiring QC analysis.	Frozen (-7°C to -15°C) or iced to 4°C (±2°C).	48 hours			
	Number 7.	Samples with MS/MSD	-	9	1	10	5g	unury 515.		48 hours			

Notes

Minimum volume/mass to be collected in order to ensure sample analysis can be performed.

Check Regional guidance regarding use of acid as a preservative of samples that may contain carbonates, residual chlorine, and other oxidants.

This technical holding time is calculated from the time of sample collection to sample extraction. Sample extracts are to be analyzed within 40 days of extraction. It is recommended that samplers ship samples to the laboratory on the same day that they are collected, or as soon as possible thereafter.

Check Regional guidance regarding use of acid preservatives when testing for carbonates, residual chlorine, and other oxidants.

Table 3-3. Sample Collection Requirements for CLP SOW SOM01 (SVOAs, Pesticides and Aroclors)

Analysis	Matrix	Containers	Minimum Volume/ Mass	Important Notes	Preservative	Technical Holding Time
Semivolatile Analytes	Water	See Table 2-2, Reference Number 5.	2L	If amber containers are not available, the samples should be protected from light.	Cool all samples to 4°C (±2°C) immediately after collection. DO NOT FREEZE water samples.	7 days
Analytes	Soil/ Sediment	See Table 2-2, Reference Numbers 3 and 4.	Fill to capacity		Cool all samples to 4°C (±2°C) immediately after collection.	14 days
Pesticides/	Water	See Table 2-2, Reference Number 5.	2L	If amber containers are not available, the samples should be protected from light.	1	7 days
Aroclors	Soil/ Sediment	See Table 2-2, Reference Numbers 3 and 4.	Fill to capacity		Cool all samples to 4°C (±2°C) immediately after collection.	14 days

Notes

Minimum volume/mass to be collected in order to ensure sample analysis can be performed. Check Regional guidance regarding use of acid as a preservative of samples that may contain carbonates, residual chlorine, and other oxidants.

This technical holding time is calculated from the time of sample collection to sample extraction. Sample extracts are to be analyzed within 40 days of extraction. It is recommended that samplers ship samples to the laboratory on the same day that they are collected, or as soon as possible thereafter.

⁴ Check Regional guidance regarding use of acid preservatives when testing for carbonates, residual chlorine, and other oxidants.

Table 3-4. Sample Collection Requirements for CLP SOW ILM05

Analysis	Matrix	Containers	Minimum Volume/ Mass ¹	Important Notes	Preservative	Technical Holding Time ⁴
Metals/ICP- AES and/or Mercury by CVAA	Water	See Table 2-2, Reference Number 2.	1L	If collecting for both ICP-AES AND ICP-MS methods, a separate 1L volume of sample must be collected for each method per sample location.	Acidify to pH < 2 with HNO ₃ and cool to 4°C (±2°C) immediately after collection. ² NOT FREEZE water samples.	6 months for all metals except Mercury (28 days)
	Soil/ Sediment	See Table 2-2, Reference Number 3.	Fill to capacity		Cool to 4°C (±2°C) immediately after collection.	6 months
Cyanide/ Spectrophoto metric Determination	Water	See Table 2-2, Reference Number 2.	1L		To neutralize residual chlorine, immediately upon collection, add 0.6 g ascorbic acid for each liter of sample collected. Add NaOH until pH >12 and cool to 4°C (±2°C) immediately after collection. ⁵ DO NOT FREEZE water samples.	14 days
	Soil/ Sediment	See Table 2-2, Reference Number 3.	Fill to capacity		Cool to 4°C (±2°C) immediately after collection.	14 days

Notes

¹ Minimum volume/mass to be collected in order to ensure sample analysis can be performed.

² Check Regional guidance regarding use of acid as a preservative of samples that may contain carbonates, residual chlorine, and other oxidants.

Samplers must test for sulfide and oxidizing agents (e.g., chlorine) in aqueous samples in the field upon collection. Please refer to the SAP and Appendix C for guidance. Sulfides adversely affect the analytical procedure. The following can be done to test for and neutralize sulfides. Place a drop of the sample on lead acetate test paper to detect the presence of sulfides. If sulfides are present, treat 25 mL more of the sample than that required for the cyanide determination with powdered cadmium carbonate or lead carbonate. Yellow cadmium sulfide or black lead sulfide precipitates if the sample contains sulfide. Repeat this operation until a drop of the treated sample solution does not darken the lead acetate test paper. Filter the solution through a dry filter paper into a dry beaker, and from the filtrate measure the sample to be used for analysis. Avoid a large excess of cadmium carbonate and a long contact time in order to minimize a loss by complication or occlusion of cyanide on the precipitated material. Sulfide removal should be performed in the field, if practical, prior to pH adjustment with NaOH.

⁴ This technical holding time is calculated from the time of sample collection to sample extraction. Sample extracts are to be analyzed within 40 days of extraction. It is recommended that samplers ship samples to the laboratory on the same day that they are collected, or as soon as possible thereafter.

3.2 Complete Documentation

Samplers must complete all documentation, including the recording of the CLP Sample Number on the sample container or bottle, sample labels, and chain-of-custody seals (as appropriate), the completion of the TR/COC Record, and the completion of field operations records (as necessary).

Samplers should use the FORMS II Lite software to create and print sample labels and the TR/COC Record. Samplers can create and print out two copies of a sample label and attach one to the sample container or bottle, and place the other on the sample tag that may be attached to the sample container or bottle.

Samplers are expected to review their project plans to determine what documentation they are expected to include during a sampling event. It is highly recommended that samplers provide documentation, even if the Region does not require it.



Under no circumstances should the site name appear on any documentation being sent to the laboratory.

An example of a packaged sample is shown in Figure 3-1. A description of each type of documentation and instructions for accurate completion are included in the following sections.

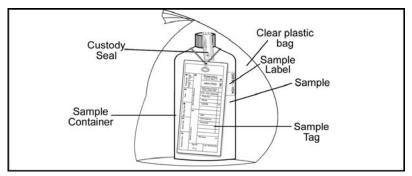


Figure 3-1. Packaged Sample with Identification and Chain-of-Custody Documentation (Excluding TR/COC Record)

3.2.1 Identify a Sample with a CLP Sample Number and SMO-assigned Case Number

The CLP Sample Number and SMO-assigned Case Number <u>must</u> be recorded on each sample taken during a sampling event (see Section 1.4.1.1). Samplers can record these numbers on the sample bottle or container using permanent ink. The numbers must also be recorded on the sample tag, if required.



Dissolved metal samples and total metal samples taken from the same sampling location cannot have the same CLP Sample Number because two different sets of data will be generated.

3.2.2 Complete TR/COC Records

A Traffic Report is used as physical evidence of sample custody and as a permanent record for each sample collected. A chain-of-custody record documents the exchange and transportation of samples from the field to the laboratory.

The ASB requires samplers to use the FORMS II Lite software to create documentation for all CLP sampling efforts. For assistance with obtaining or using the FORMS II Lite software, please contact the FORMS II Lite Help Desk at 703-818-4200 from 9:00 AM - 5:00 PM ET.

To meet CLP sample documentation and chain-of-custody requirements, the sampler must attach a separate TR/COC Record to each cooler they ship. The TR/COC Record must document each sample within the cooler. Samples shipped in other coolers should not be documented. This practice maintains the chain-of-custody for all samples in case of incorrect shipment.

If more than one TR/COC Record is used for the samples within one cooler, all of the records must have complete header information and original signatures. Samplers are responsible for the care and custody of samples from the time of collection to the time of shipment to the laboratories for analysis. A sample is considered under custody if:

- It is in possession or in view after being in possession;
- It was in possession and then secured or sealed to prevent tampering; or
- It was in possession when placed in a secured area.

Each time the custody of samples is turned over to another person, the TR/COC Record must be signed off by the former custodian and accepted by the new custodian. Samplers are, therefore, responsible for properly completing any forms or other Region-required documentation used to establish the chain-of-custody for each sample during a sampling event.

3.2.2.1 Complete a TR/COC Record Using the FORMS II Lite Software

Once the sampler inputs sample collection information into FORMS II Lite, a TR/COC Record will be generated electronically. The software automatically displays only the information to be entered by the sampler. FORMS II Lite then generates a laboratory and a Regional copy of the TR/COC Record (see Figures 3-2 through 3-5). The sampler can print out multiple copies of the TR/COC Record as necessary. The sampler must sign and submit original copies of the TR/COC Record as appropriate.

An electronic TR/COC Record created using the FORMS II Lite software contains basic header information; however, the sampler can also include some additional detailed information. For example, not only is the sample matrix listed on the electronic TR/COC Record, but the name of the sampler taking the sample can also be entered. Samplers should note that certain information will not appear on the electronic TR/COC Record (e.g., matrix and preservative descriptions).

3.2.2.2 Indicate Modified Analysis on FORMS II Lite TR/COC Records

When completing a TR/COC Record using FORMS II Lite, the sampler should identify any samples that will be analyzed using a CLP Modified Analysis. Samplers should indicate use of a Modified Analysis by creating a new analysis within the FORMS II Lite Wizard or through the FORMS II Lite Reference Tables. This newly-created analysis should contain the Modification Reference Number within the name assigned to the analysis. For example, if a Region submits a Modified Analysis for an additional analyte, and SMO assigns the Modification Reference Number 1301.0, the FORMS II Lite analysis could be named "VOA by M.A. 1301.0". The associated abbreviation for this analysis could be "VOA M.A.". If you have any questions regarding identification of Modified Analysis using FORMS II Lite, please contact the FORMS II Lite Help Desk at 703-818-4200 from 9:00 AM - 5:00 PM ET.

3.2.2.3 Make Manual Edits to Printed FORMS II Lite TR/COC Records

If a FORMS II Lite TR/COC Record has been printed and deletions or edits need to be made by the sampler, the following procedures must be followed:

- If making a deletion, manually cross out the information to be disregarded from the TR/COC Record, initial and date the deletion.
- If making an addition, enter the new information and initialsign and date the newly added information.



All modifications made on a printed TR/COC Record must be initialed and dated.

Date Shipped:	2/20/2001		Chain of Custody I	Record	Sampler Signature:		For Lab Use Only				
Carrier Name:	DHL		Relinquished By	(Date / Time)	Received By	(Date / Time)	Lab Con	ntract No:			
Airbill: Shipped to:	121212 Organic Laboratory	,	1				Unit Prid	ce:			
отррес то.	1234 Smith Drive		2				Transfer	15			
	Anywhere, USA 12 (123) 456-7890	345	3					ntract No:			
			4				Unit Pric				
ORGANIC SAMPLE No.	MATRIX/ SAMPLER	CONC/ TYPE	ANALYSIS/ TURNAROUND	TAG No./ PRESERVATIVE/ Bottles	STATION LOCATION	SAMPLE COLI DATE/TIMI	ECT	INORGANIC SAMPLE No.	FOR LAB USE ONLY Sample Condition On Receipt		
C0075	Industrial Process Wastewater/ BOBBY	H/C	BNA/PEST (21), VOA 6 (21)	3486, 6487 (2)	LOCATION ONE	S: 2/20/2001 E: 2/23/2001	16:02 16:02	MC0075			
C0076	SAMPLER Ground Water/ JOE SAMPLER	L/C	BNA/PEST (21), VOA 6 (21)	6494, 6495 (2)	LOCATION TWO	S: 2/20/2001 E: 2/21/2001	16:01 16:01	MC0076			
C0077	Industrial Effluent Wastewater/ JOE SAMPLER	t M/G	BNA/PEST (21), VOA 6 (21)	3502, 6503 (2)	LOCATION ONE	S: 2/16/2001 E: 2/20/2001	15:55 15:55	MC0077			
nipment for Case omplete ?N	Sample(s) to C0077	be used	for laboratory QC:	Additional Sample	er Signature(s):	Cooler Tempera Upon Receipt:	ture	Chain of Custody	Seal Number:		
						(33)					
nalysis Key:	Concentrati	on: L=	Low, M = Low/Medium, H =	High Type/D	esignate: Composite = C, G	rab = G		Custody Seal Inta	ct? _ Shipment Iced? _		

Figure 3-2. Organic Traffic Report & Chain of Custody Record (Laboratory Copy)

Date Shipped:	2/20/2001	The state of the s	Chain of Custody	Record	Sampler		For La	b Use Only	
Carrier Name:	DHL		Relinguished By	(Date / Time)	Signature: Received By (Date / Time)			ntract No:	
Airbill:	121212		1				200 100000		
Shipped to:	Inorganic Laboratory 1234 Smith Drive		2				Unit Pri	38	
	Anywhere, USA 12345 (123) 456-7890		3				Transfer		
	(123) 430-7030		1870				Lab Cor	ntract No:	
100 No. Service Louise 17 (44 Co.	100 000 00 To 100		4			************	Unit Pri	ce:	AND ATTICLE CONTROL OF THE CONTROL O
INORGANIC SAMPLE No.		CONC/ TYPE	ANALYSIS/ TURNAROUND	TAG No./ PRESERVATIVE/ Bottles	STATION LOCATION	SAMPLE COL DATE/TIM		ORGANIC SAMPLE No.	FOR LAB USE ONLY Sample Condition On Receipt
MC0075	Industrial Process Wastewater/ BOBBY	H/C	Al (21), Ba (21), Ca (21), Cr (21), TM/CN (21)	6481, 6482, 6483, 6484, 6485 (5)	LOCATION ONE	S: 2/20/2001 E: 2/23/2001	16:02 16:02	C0075	
MC0076	SAMPLER Ground Water/ JOE SAMPLER	L/C	Al (21), Ba (21), Ca (21), Cr (21), TM/CN (21)	6489, 6490, 6491, 6492, 6493 (5)	LOCATION TWO	S: 2/20/2001 E: 2/21/2001	16:01 16:01	C0076	
MC0077	Industrial Effluent Wastewater/ JOE SAMPLER	M/G		6497, 6498, 6499, 6500, 6501 (5)	LOCATION ONE	S: 2/16/2001 E: 2/20/2001	15:55 15:55	C0077	
ihipment for Case	Sample(s) to b	e used f	for laboratory QC:	Additional Sample	er Signature(s):	Cooler Tempera	thure.	Chain of Custody	r Saal Number:
omplete?N	MC0077					Upon Receipt:			0007 (1983) 1000 (4 A X (64 PM) (4 M))
			Low, M = Low/Medium, H	High Type/De	esignate: Composite = C, G	rab = G		Custody Seal Inta	sct? Shipment Iced?

Figure 3-3. Inorganic Traffic Report & Chain of Custody Record (Laboratory Copy)

Region: Project Code:	3 QW-123			Date Shipped: Carrier Name:	2/20/2001 DHL		Chai	in of Custody	Record		Sampler Signature:	
Account Code: CERCLIS ID:	ACCT000			Airbill:	121212		Relin	quished By	(Date	/ Time)	Received By	(Date / Time)
Spill ID:	ID3			Shipped to:	Organic Laborato		1				9	
Site Name/State					Anywhere, USA 1 (123) 456-7890		2					
Project Leader: Action:	DAN SAMP Other	.EK			(120) 400-1000		3				8	
Sampling Co:	SMITH CO.						4					
ORGANIC SAMPLE No.	MATRIX/ SAMPLER	CONC/ TYPE	ANALYSIS/ TURNAROUND	TAG N PRESERVATI		STATION			COLLECT E/TIME		GANIC PLE No.	QC Type
	Industrial Process Wastewater/ BOBBY	H/C	BNA/PEST (21), VOA (21)	6486, 6487 (2)		LOCATION OF	NE	S: 2/20/2001 E: 2/23/2001	16:02 16:02	MC0075		-
	SAMPLER Ground Water/	L/C	BNA/PEST (21), VOA	6494, 6495 (2)		LOCATION TV	VO	S: 2/20/2001	16:01	MC0076		Spike
00077	JOE SAMPLER Industrial	M/G	(21) BNA/PEST (21), VOA	6502 6502 (2)		LOCATION OF		E: 2/21/2001 S: 2/16/2001	16:01 15:55	MC0077		_
	Effluent Wastewater/ JOE SAMPLER		(21)	0002, 0000 (2)		LOGATION OF		E: 2/20/2001	15:55	Mooori		
Shipment for Case	I Samula(a		and for laboratory OC:		Additional Sample	ur Sinnatura(e)				Į,	Chain of Custody :	Seal Number
Complete? N	C0077	to be us	sed for laboratory QC:		Additional Sample	er Signature(s).					Shain of Gustody s	Sear Number:
Analysis Key: BNA/PEST = CL	Concenti P TCL Semivolatil		L = Low, M = Low/Medium Pesticides/PC, VOA = CL		Type/Designate:	Composite = C	, Grab	= G			Shipment Iced?	

Figure 3-4. Organic Traffic Report & Chain of Custody Record (Region Copy)

Region: Project Code:	3 QW-123			Date Shipped: Carrier Name:	2/20/2001 DHL		Cha	in of Custody R	ecord		Sampler Signature:	
Account Code: CERCLIS ID:	ACCT000			Airbill:	121212	- 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100	Relin	quished By	(Date	/ Time)	Received By	(Date / Time)
Spill ID:	ID3			Shipped to:	Clayton Envi	Inc	2					
Site Name/State Project Leader:					22345 Roeth Novi MI 483 (248) 344-17	75	3					
Action: Sampling Co:	Other SMITH Co	0			(240) 344-17	70	4					
INORGANIC SAMPLE No.	MATRIX/ SAMPLER	CONC/ TYPE	ANALYSIS/ TURNAROUND	TAG PRESERVAT		STATION LOCATION	_	SAMPLE C			L GANIC PLE No.	QC Type
AC0075	Industrial Process Wastewater/ BOBBY SAMPLER	H/C	Al (21), Ba (21), Ca (21), Cr (21), TM/CN (21)	6481, 6482, 64 6485 (5)	83, 6484,	LOCATION OF	NE		16:02 16:02	C0075		3.75
MC0076	Ground Water/ JOE SAMPLER	L/C	Al (21), Ba (21), Ca (21), Cr (21), TM/CN		91, 6492,	LOCATION TV	VO		16:01 16:01	C0076		Spike
1C0077	Industrial Effluent Wastewater/ JOE SAMPLER	M/G	(21) AI (21), Ba (21), Ca (21), Cr (21), TM/CN (21)		99, 6500,	LOCATION OF	NE		15:55 15:55	C0077		-
hipment for Case complete? N	Sample	e(s) to be us	sed for laboratory QC:		Additional Sa	impler Signature(s):					Chain of Custody	Seal Number:
Analysis Key: Al = Aluminum,	20 M		L = Low, M = Low/Medium , Cr = Chromium, TM/C		Type/Design		, Grab	= G		Shipment Iced?		

Figure 3-5. Inorganic Traffic Report & Chain of Custody Record (Region Copy)

3.2.3 Complete and Attach Custody Seals

Custody seals are usually pre-printed stickers that are signed (or initialed) and dated by the sampler after sample collection and placed on sample bottles or containers and/or shipping coolers or containers (see Figure 3-6). The custody seals document who sealed the sample container and verifies that the sample has not been tampered with. The seals must be placed such that they will break if the sample bottle or container or the shipping cooler or container is tampered with or opened after leaving custody of samplers. Custody seals can also be used to maintain custody of other items such as envelopes containing videotapes of the sample collection process.



Custody seals should never be placed directly onto a coring tool used as a transport device (e.g., 5 g Sampler) or tared, 40 mL closed-system vials. The seals must be placed on the bag for the coring tool used as a transport device, or on the bag used to enclose the vials. Refer to Appendix B for details.

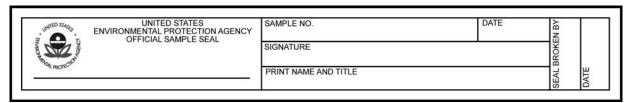


Figure 3-6. Custody Seal

Instructions for completing and attaching a custody seal are included in Table 3-5.

Step	Action	Important Notes
1	Record the CLP Sample Number.	The space for the CLP Sample Number does not need to be completed on custody seals being placed on the opening of a cooler, only on those being placed on the opening of sample bottles or containers.
2	Record the month, day, and year of sample collection.	
3	Sign the seal in the Signature field.	
4	Print your name and title in the Print Name and Title field.	
5	Place the custody seal over the edge of the sample bottle or container such that it will break if tampered with.	Custody seals can be placed directly on any sample container except for coring tools used as a transport device (e.g., 5 g Samplers) and tared VOA bottles. If packing coring tools used as a transport device or tared VOA bottles, place them in a clear plastic bag and place the custody seal on the outside of the bag.
6	If possible, cover the custody seal with clear plastic tape to protect it.	Take special care to not place the protective tape over the seal in such a way that it can be removed and then re-attached without signs of tampering.

Table 3-5. Completing and Attaching a Custody Seal

The use and type of custody seals can vary by Region or collecting organization. Samplers should obtain the appropriate custody seals and specific instructions for correctly attaching them from the RSCC.

3.2.4 Complete and Attach Sample Labels

Samplers affix sample labels to each sample container. A sample label must contain the associated CLP Sample Number (either written or pre-printed), SMO-assigned Case Number, and the preservative used. It must also denote the analysis/fraction. Samplers may also include additional information such as the station location or the date/time of collection. Samplers should use FORMS II Lite to create and print sample labels. The sampler can print two labels and attach one to the sample container or bottle, and place the other label on the sample tag that should also be attached to the sample container or bottle. The

labels should then be covered with clear packaging tape to protect the label and maintain legibility. If handwriting a sample label, the sampler should complete the label information using waterproof ink, place the label on the outside of the sample bottle or container, then cover the label with clear packaging tape to protect the label and maintain legibility (see Figure 3-1).



Do not attach labels to tared VOA sample vials. A label should already be pre-attached to the tared vial.

3.2.5 Complete and Attach Sample Tags

To support use of sample data in potential enforcement actions, sample characteristics other than on-site measurements (e.g., pH, temperature, conductivity) can be identified with a sample tag. Typically, site-specific information is written on the tags using waterproof ink. The use and type of sample tags may vary by Region. For each sampling event, samplers should receive the required sample tags and type of information to include from the RSCC. The sampler can use FORMS II Lite to create and print out multiple sample labels, one of which can be attached to the sample tag and then covered with clear packaging tape to protect the label and maintain legibility. If FORMS II Lite-created sample labels are not available, a detailed set of instructions for completing and attaching a handwritten sample tag are included in Table 3-6.



The use and type of sample tags may vary among Regions.

Table 3-6. Completing and Attaching a Handwritten Sample Tag

Step	Action	Important Notes
1	Under the "Remarks" heading, record the CLP Sample Number and SMO-assigned Case Number.	Make sure to record the correct CLP Sample Number and SMO-assigned Case Number in a legible manner.
2	Record the project code (e.g., Contract Number, Work Assignment Number, Interagency Agreement Number, etc.) assigned by USEPA.	
3	Enter the station number assigned by the sampling team coordinator.	
4	Record the month, day, and year of sample collection.	
5	Enter the military time of sample collection (e.g., 13:01 for 1:01 PM).	
6	Identify the designate and place an "X" in either the "Comp." or "Grab" box if the sample is either a composite or grab sample.	
7	Record the station location.	
8	Sign the sample tag in the Signature area.	
9	Place an "X" in the box next to Yes or No to indicate if a preservative was added to the sample.	
10	Under "Analyses", place an "X" in the box next to the parameters for which the sample is to be analyzed.	
11	Leave the box for "Laboratory Sample Number" blank.	
12	It is recommended that the sample tag be attached to the neck of the sample bottle or container using regular string, stretch string, or wire (see Figure 3-1).	Do NOT use wire to attach a sample tag to a metals sample.

Station No. 3 Mo./Day/Year 4 5 2 Designate: 6 Time **Project Code** 01/10/2004 8:45 AM Grab 1 Comp. 00-030 Sampler's (Signature) Station Location 7 8 John Smith 3-3001 Tag COD, BOD Anions Solids (TSS) (TDS) (SS) Preservative: SVOA organics ABN Remarks Pesticides VOA organics Metals Cyanide Phenolics ANALYSES TOC, Nutrients DD001 Lab. Sample No 9 1 10 ×

An example of a completed sample tag is included in Figure 3-7 below:

Figure 3-7. Completed Sample Tag

3.3 Provide Sample Receipt

After samples have been taken from private property, the sampler should prepare a receipt for these samples and provide this receipt to the property owner. This is especially important when sampling on private property since these samples could be used during future litigation and the receipt will verify that the owner granted approval for the removal of the samples from the property. An example of a sample receipt created using FORMS II Lite is shown in Figure 3-8.

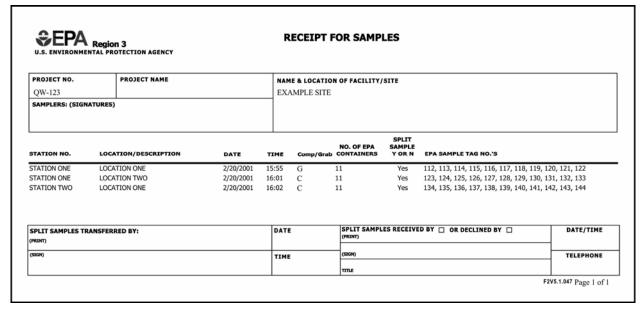


Figure 3-8. Sample Receipt Created Using the FORMS II Lite Software

3.4 Pack and Ship Samples

Once the samples have been collected, it is very important that the sampler properly package the samples for shipment and ensure that the samples are sent to the appropriate laboratory as quickly as possible. Prompt and proper packaging of samples will:

- Protect the integrity of samples from changes in composition or concentration caused by bacterial growth or degradation from increased temperatures;
- Reduce the chance of leaking or breaking of sample containers that would result in loss of sample volume, loss of sample integrity, and exposure of personnel to toxic substances; and
- Help ensure compliance with shipping regulations.

3.4.1 Sample Containers

Once samples are collected, they must be stored in conditions that maintain sample integrity. All samples should be placed in shipping containers or other suitable containers with ice to reduce the temperature as soon as possible after collection. Ideally, all samples should be shipped the day of collection for overnight delivery to the laboratory. If samples cannot be shipped on the day of collection, the sample temperature should be maintained at 4° C ($\pm 2^{\circ}$ C) until they are shipped to the laboratory.

One CLP RAS sample may be contained in several bottles and vials. For example, one soil sample may consist of all containers needed for three of the analytical fractions available under this service (i.e., SVOA fraction, Pesticide fraction, and Aroclor fraction), even though the fractions are collected in separate containers. Therefore, the analysis to be performed and the matrix type will determine the type of container(s) that will be used, as well as the volume that must be collected for that particular sample fraction.

3.4.2 Inventory of Samples and Documentation

Prior to shipment, samplers should conduct an inventory of the contents of the shipping cooler or container against the corresponding TR/COC Record when packing for shipment to laboratories. An inventory will ensure that the proper number of containers have been collected for each analysis of the samples, that the required PE and QC samples and cooler temperature blanks are included, and the correct Sample Numbers and fractions have been assigned to each sample.

3.4.3 Shipping Regulations

Sample shipping personnel are legally responsible for ensuring that the sample shipment will comply with all applicable shipping regulations. For example, hazardous material samples must be packaged, labeled, and shipped in compliance with all IATA Dangerous Goods regulations or DOT regulations and USEPA guidelines. Refer to Appendix B for detailed shipping guidelines when using SW-846 Method 5035A to preserve and ship samples.

3.4.4 Sample Packaging for Shipment

Samplers are responsible for the proper packaging of samples for shipment. To ensure that samples are appropriately packaged (e.g., to avoid breakage and/or contamination) the sampler should consult their respective project plans to determine the proper packing and shipping procedures. The sampler must determine the sample type, pack the shipping containers correctly, include necessary paperwork, label and seal shipping containers or coolers, and ship the samples.

3.4.4.1 Determine the Sample Type and Container

Samplers should know what kinds of samples they are handling to ensure proper packaging. Samplers should refer to their appropriate project plans to determine which type of sample container should be used for each type of sample being taken during the sampling event.



Please follow Regional guidance with reference to samples containing dioxin or radioactive waste.

3.4.4.2 Pack Shipping Containers

It is imperative that samples are correctly and carefully packed in shipping containers to prevent the sample containers from breaking or leaking. Samplers must prepare and pack a shipping cooler or container according to the instructions outlined in Table 3-7.

Table 3-7. Packing Samples for Shipment

Step	Action	Important Notes
1	Seal all drain holes in the shipping container, both inside and out, to prevent leakage in the event of sample breakage.	
2	Check all lids/caps to make sure the samples are tightly sealed and will not leak.	
3	Seal samples within a clear plastic bag.	Custody seals can be placed directly on any sample container except for coring tools used as a transport device (e.g., 5 g Samplers) and tared VOA bottles. If packing coring tools used as a transport device or tared VOA bottles, place them in a clear plastic bag and place the custody seal on the outside of the bag.
4	Fully chill samples to 4°C (±2°C) prior to placement within suitable packing materials.	
5	Prior to placing samples within the shipping cooler, it is recommended that samplers line shipping containers with non-combustible, absorbent packing material.	
6	Place samples in CLEAN, sealed, watertight shipping containers (metal or hard plastic coolers).	
7	Conduct an inventory of the contents of the shipping cooler/container against the corresponding TR/COC Record.	
8	Cover samples in double-bagged ice to prevent water damage to packing materials.	Do NOT pour loose ice directly into the sample cooler. The ice is used to maintain the temperature of the samples within the shipping cooler.
9	It is recommended a temperature blank be included within each cooler being shipped.	The temperature blank is generally a 40 L vial filled with water and labeled "temperature blank" but does not have a Sample Number.
10	Ensure that the site name or other site-identifying information does not appear on any documentation being sent to the laboratory.	The laboratory should not receive any site-identifying information.

3.4.4.3 Include Necessary Paperwork

Samplers must properly place the necessary paperwork in the shipping cooler. All paperwork must be placed in a plastic bag or pouch and then secured to the underside of the shipping cooler lids (see Figure 3-9).

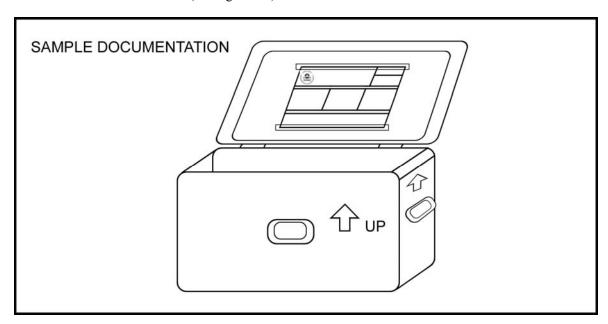


Figure 3-9. Sample Cooler with Attached TR/COC Record and Cooler Return Documentation

Necessary paperwork includes TR/COC Records and sample weight logs (see Figure 3-10), if required (for VOA samples). Samplers should contact their RSCC (or designee) for specific paperwork requirements.

Shinned to:	AAA Testing L	abaratan (mple Wei	9 9		Case No.	39563	
	1700 Mill Aver						DAS No.		
	Houston TX 7							DAS34	
	(281) 983-123	4					Date Shipped:	9/29/2003	
Sample No.	Matrix	Analysis	Preservative	Bottle/ Tag Number	Tared Weight (g)	Final Weight (g)	Sample Weight (g)	Laboratory Weight	Traffic Report No.
C0036	Subsurface Soil (>12")	CLP TCL Volatiles	Ice Only	199548	32.80	37.20	4.40		3-103018225-092903-000
C0036	Subsurface Soil (>12")	CLP TCL Volatiles	Ice Only	199547	32.10	38.30	6.20		3-103018225-092903-000
C0036	Subsurface Soil (>12")	CLP TCL Volatiles	Ice Only	199549	31.20	38.60	7.40		3-103018225-092903-000
C0037	Surface Soil (0"-12")	CLP TCL Volatiles	Ice Only	199552	32.00	36.90	4.90		3-103018225-092903-000
C0037	Surface Soil (0"-12")	CLP TCL Volatiles	Ice Only	199551	32.40	37.10	4.70		3-103018225-092903-000
C0037	Surface Soil (0"-12")	CLP TCL Volatiles	Ice Only	199550	31.90	35.90	4.00		3-103018225-092903-000
Completed By	y:			Da	te:				

Figure 3-10. Sample Weight Log

3.4.4.4 Return Sample Shipping Coolers

CLP laboratories must routinely return sample shipping coolers within 14 calendar days following shipment receipt. Therefore, the sampler should also include cooler return instructions with each shipment. The sampler (not the CLP laboratory) is responsible for paying for return of the cooler and should also include shipping airbills bearing the sampler's account number, as well as a return address to allow for cooler return.

3.4.4.5 Label and Seal Sample Shipping Coolers

After samples are packaged within shipping coolers, samplers must carefully secure the top and bottom of the coolers with tape, place return address labels clearly on the outside of the cooler, and attach the required chain-of-custody seals (see Figure 3-11).

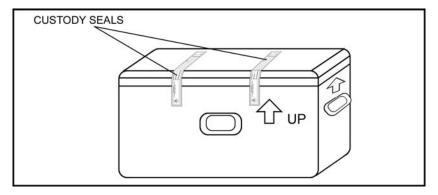


Figure 3-11. Shipping Cooler with Custody Seals

If more than one cooler is being delivered to a laboratory, samplers should mark each cooler as "1 of 2", "2 of 2", etc. In addition, samplers must accurately complete and attach shipping airbill paperwork for shipment of the samples to the laboratory. An airbill, addressed to the Sample Custodian of the receiving laboratory, should be completed for each cooler shipped. Samplers should receive the correct name, address, and telephone number of the laboratory to which they must ship samples from the RSCC or SMO. To avoid delays in analytical testing, samplers should make sure they are sending the correct types of samples to the correct laboratory when collecting samples for multiple types of analysis. For example, inorganic samples may be shipped to one laboratory for analysis, while organic samples may need to be shipped to another laboratory.

3.4.4.6 Ship Samples

The sampling contractor should ensure that samplers know the shipping company's name, address, and telephone number. In addition, they should be aware of the shipping company's hours of operation, shipping schedule, and pick-up/drop-off requirements.

Overnight Delivery

It is imperative that samples be sent via overnight delivery. Delays caused by longer shipment times may cause technical holding times to expire, which in turn may destroy sample integrity or require the recollection of samples for analysis.

Saturday Delivery

For shipping samples for Saturday delivery, the sampler MUST contact the RSCC (or their designee) or SMO so that SMO will receive the delivery information by 3:00 PM ET on the Friday prior to delivery.

3.4.5 Shipment Notification

When samples are shipped to CLP Laboratories, samplers <u>must immediately</u> report all sample shipments to the RSCC (or their designee) or to SMO. **Under no circumstances should the sampler contact the laboratory directly.** If samplers are shipping samples after 5:00 PM ET, they must notify the RSCC (or

designee) or SMO by 8:00 AM ET on the following business day. Samplers should receive the name and phone number of the appropriate SMO coordinator to contact from the Region/RSCC.

Samplers must provide the following information to the RSCC (or their designee) or to SMO:

- Name and phone number at which they can easily be reached (preferably closest on-site phone number if still in the field);
- SMO-assigned Case Number (see Section 2.4.1);
- Number, concentration, matrix and analysis of samples being shipped;
- Name of laboratory (or laboratories) to which the samples were shipped;
- Airbill number(s);
- Date of shipment;
- Case status (i.e., whether or not the Case is complete);
- Problems encountered, special comments, or any unanticipated issues;
- When to expect the next anticipated shipment; and
- An electronic export of the TR/COC Record (must be sent as soon as possible after sample shipment). For information regarding electronic export of TR/COC Records, refer to the following Web site:

http://www.epa.gov/superfund/programs/clp/f2lsubmit.htm



For Saturday delivery, samplers MUST contact the RSCC (or their designee) or SMO so that SMO will receive the delivery information by 3:00 PM ET on the Friday prior to delivery.

Samplers should be aware if their Region requires them to notify the RSCC (or designee) and/or SMO of sample shipment.

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Appendix A: Functions within a Sampling Project

The following table describes Quality Assurance Project Plan (QAPP) requirements taken from *EPA Requirements for Quality Assurance Project Plans* (EPA QA/R-5).

Functions Within a Sampling Project	Elements of that Function				
Project Management					
Project/Task Organization	Identifies the individuals or organizations participating in the project and defines their specific roles and responsibilities.				
Problem Definition/Background	States the specific problem to be solved or decision to be made and includes sufficient background information to provide a historical and scientific perspective for each particular project.				
Project/Task Description	Describes the work to be performed and the schedule for implementation to include: • Measurements to be made during the course of the project; • Applicable technical, regulatory, or program-specific quality standards, criteria, or objectives; • Any special personnel and equipment requirements; assessment tools needed; and • A work schedule and any required project and quality records, including types of reports needed.				
Quality Objectives and Criteria	Describes the project quality objectives and measurement performance criteria.				
Special Training/Certification	Ensures that any specialized training for non-routine field sampling techniques, field analyses, laboratory analyses, or data validation should be specified.				
Documents and Records	 Itemizes the information and records that must be included in the data report package and specifies the desired reporting format for hard copy and electronic forms, when used. Identifies any other records and/or documents applicable to the project such as audit reports, interim progress reports, and final reports that will be produced. Specifies or references all applicable requirements for the final disposition of records and documents, including location and length of retention period. 				
	Data Generation and Acquisition				
Sampling Process Design (Experimental Design)	 Describes the experimental design or data collection design for the project. Classifies all measurements as critical or non-critical. 				
Sampling Methods	 Describes the procedures for collecting samples and identifies sampling methods and equipment. Includes any implementation requirements, support facilities, sample preservation requirements, and materials needed. Describes the process for preparing and decontaminating sampling equipment to include the disposal of decontamination by-products, selection and preparation of sample containers, sample volumes, preservation methods, and maximum holding times for sampling, preparation, and/or analysis. Describes specific performance requirements for the method. Addresses what to do when a failure in sampling occurs, who is responsible for corrective action, and how the effectiveness of the corrective action shall be determined and documented 				
Sample Handling and Custody	Describes the requirements and provisions for sample handling and custody in the field, laboratory, and transport, taking into account the nature of the samples, the maximum allowable sample holding times before extraction and analysis, and the available shipping options and schedules. Includes examples of sample labels, custody forms, and sample custody logs.				

	• Identifies the analytical methods and equipment required, including sub-sampling or extraction methods, waste disposal requirements (if any), and specific method performance
	requirements.
	• Identifies analytical methods by number, date, and regulatory citation (as appropriate). If a method allows the user to select from various options, the method citations should state exactly which options are being selected.
Analytical Methods	Addresses what to do when a failure in the analytical system occurs, who is responsible for corrective action, and how the effectiveness of the corrective action shall be determined and documented.
	Specifies the laboratory turnaround time needed, if important to the project schedule.
	Specifies whether a field sampling and/or laboratory analysis Case Narrative is required to provide a complete description of any difficulties encountered during sampling or analysis.
	Identifies required measurement QC checks for both the field and laboratory.
	• States the frequency of analysis for each type of QC check, and the spike compounds sources and levels.
Quality Control (QC)	States or references the required control limits for each QC check and corrective action required when control limits are exceeded and how the effectiveness of the corrective action shall be determined and documented.
	Describes or references the procedures to be used to calculate each of the QC statistics.
	Describes how inspections and acceptance testing of environmental sampling and measurement systems and their components will be performed and documented. Identifies and discusses the procedure by which final acceptance will be performed by independent personnel.
Instrument/Equipment Testing,	Describes how deficiencies are to be resolved and when re-inspection will be performed.
Inspection, and Maintenance	Describes or references how periodic preventative and corrective maintenance of measurement or test equipment shall be performed.
	Identifies the equipment and/or system requiring periodic maintenance.
	Discusses how the availability of spare parts identified in the operating guidance and/or design specifications of the systems will be assured and maintained.
	Identifies all tools, gauges, instruments, and other sampling, measuring, and test equipment used for data collection activities affecting quality that must be controlled, and at specific times, calibrated to maintain performance within specified limits.
Instrument/Equipment	Identifies the certified equipment and/or standards used for calibration.
Calibration and Frequency	• Describes or references how calibration will be conducted using certified equipment and/or standards with known valid relationships to nationally recognized performance standards. If no such standards exist, documents the basis for calibration.
	Indicates how records of calibration shall be maintained and traced to the instrument.
Inspection/Acceptance of Supplies and Consumables	Describes how and by whom supplies and consumables shall be inspected and accepted for use in the project.
Supplies and Consumables	States acceptance criteria for such supplies and consumables.
	Identifies any types of data needed for project implementation or decision-making that are obtained from non-measurement sources (e.g., computer databases, programs, literature files, historical databases).
Non-direct Measurements	Describes the intended use of data.
	Defines the acceptance criteria for the use of such data in the project. Consider the state of the
	Specifies any limitations on the use of the data.
Data Management	• Describes the project data management scheme, tracing the data path from generation in the field or laboratory to their final use or storage.
, , , , , , , , , , , , , , , , , , ,	Describes or references the standard record-keeping procedures, document control system, and the approach used for data storage and retrieval on electronic media.

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Appendix B: CLP Sample Collection Guidelines for VOAs in Soil by SW-846 Method 5035A

A. Preferred Options for the Contract Laboratory Program (CLP) are Options 1, 2, and 3:



Soil samples must be placed on their sides prior to being frozen.

Option 1.

Closed-system Vials:

Container - tared or preweighed 40 mL VOA Vials containing a magnetic stir bar.

Collect 5 g of soil per vial (iced or frozen in the field).

Regular Samples 3 Vials - Dry (5 g soil per vial)

1 Vial - Dry (filled with soil, no headspace)

4 Total Vials

Regular Samples 9 Vials - Dry (5 g soil per vial)

Requiring QC Analysis 1 Vial - Dry (filled with soil, no headspace)

10 Total Vials

Option 2.

Closed-system Vials Containing Water:

Container - tared or pre-weighed 40 mL VOA vials containing a magnetic stir bar and 5 mL water.

Collect 5 g of soil per vial (iced or frozen in the field).

Regular Samples 2 Vials with water added (5 g soil and 5 mL water per vial)

1 Vial - Dry (5 g soil in vial)

1 Vial - Dry (filled with soil, no headspace) 4 Total Vials (2 with water and 2 dry)

Regular Samples 6 Vials with water added (5 g soil and 5 mL water per vial)

Requiring QC Analysis 5 Vials - Dry (5 g soil per vial)

1 Vial - Dry (filled with soil, no headspace)
12 Total Vials (6 with water and 6 dry)

Option 3.

Coring Tool used as a Transport Device

Container - 5 g Samplers or equivalent.



All Samplers should be iced or frozen in the field and bagged individually.

Regular Samples 3 Samplers (5 g soil per Sampler)

1 Vial - Dry (filled with soil, no headspace)

4 Total (3 Samplers and 1 Vial)

Regular Samples 9 Samplers (5 g soil per Sampler)

Requiring QC Analysis 1 Vial - Dry (filled with soil, no headspace)

10 Total (11 Samplers and 1 Vial)

B. Options 4, 5, and 6 are NOT preferred options for the CLP:

Option 4.

Closed-system Vials:

Container - tared or preweighed 40 mL VOA Vials containing a magnetic stir bar and preservative.

Collect 5 g of soil per vial and add Sodium bisulfate (NaHSO₄) preservative (5 mL water + 1 g NaHSO₄) - iced or frozen in the field.

Caution: This option is NOT a Preferred Option for the CLP because:

NaHSO4 preservation creates low pH conditions that will cause the destruction of certain CLP target analytes (e.g., vinyl chloride, trichloroethene, trichlorofluoromethane, cis- and trans-1,3-dichloropropene). Projects requiring the quantitation of these analytes should consider alternative sample preservation methods. NaHSO4 also cannot be used on carbonaceous soils. Check the soil before using this method of collection! Soil can be checked by placing a test sample in a clean vial, then adding several drops of NaHSO4 solution. If the soil bubbles, use Option 4b and note this issue on the TR/COC Record.

Option 4a. Samples preserved in the field

Regular Samples 2 Vials with NaHSO₄ preservative added (5g soil per vial)

1 Vial without NaHSO₄ preservative added (5g soil per vial)

1 Vial - Dry (filled with soil, no headspace)

4 Total Vials (2 with NaHSO₄ preservative and 2 without)

Regular Samples4 Vials with NaHSO₄ preservative added (5g soil per vial) **Requiring QC Analyses**5 Vials without NaHSO₄ preservative added (5g soil per vial)

1 Vial - Dry (filled with soil, no headspace)

10 Total Vials (4 with NaHSO₄ and 6 without)

Option 4b. Samples are preserved by the laboratory (No NaHSO₄ preservative is added to these samples in the

field).

Regular Samples 3 Vials - Dry (5 g soil per vial)

1 Vial - Dry (filled with soil, no headspace)

4 Total Vials

Regular Samples 9 Vials - Dry (5 g soil per vial)

Requiring QC Analyses 1 Vial - Dry (filled with soil, no headspace)

10 Total Vials

Option 5.

Methanol Preservation (medium-level analysis only):

Container - tared or pre-weighed 40 mL VOA vials containing 5-10 mL methanol.

Collect 5 g of soil per vial (iced in the field).

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Caution: This is NOT a preferred option for the CLP because:

Samples preserved with methanol can only be analyzed by the medium-level method. Lowlevel Contract Required Quantitation Limit (CRQLs) cannot be achieved when samples are preserved this way.

Additional problems associated with use of methanol as a preservative in the field include:

- Possible contamination of the methanol by sampling-related activities (e.g., absorption of diesel fumes from sampling equipment);
- Leakage of methanol from the sample vials during shipping, resulting in loss of VOAs prior to analysis.

Regular Samples 2 Vials (5 g soil and 5-10 mL methanol per vial)

1 Vial - Dry (filled with soil, no headspace)

3 Total Vials (2 with methanol and 1 dry)

Regular Samples 6 Vials (5 g soil and 5-10 mL methanol per vial) **Requiring QC Analysis** 1 Vial -Dry (filled with soil, no headspace)

7 Total Vials (6 with methanol and 1 dry)



If shipping samples containing methanol as a preservative, a shipping label must be used to indicate methanol. This label must also contain the United Nations (UN) identification number for methanol (UN 1230), and indicate Limited Quantity.

Option 6.

Glass Containers filled with sample - No Headspace:

Container - 4 oz Glass Jars.

Glass container filled with soil with no headspace and iced.

Caution: This is NOT a preferred option for the CLP because:

Samples collected in this manner lose most of their volatile analytes prior to analysis when the sample containers are opened and sub-sampled in the laboratory. This option is only available due to Regional requirements.

Regular Samples 2 Glass Jars (4 oz) filled with sample, no headspace

1 Vial - Dry (filled with soil, no headspace)

3 Total Containers

Regular Samples 2 Glass Jars (4 oz) filled with sample, no headspace **Requiring QC Analysis**

1 Vial - Dry (filled with soil, no headspace)

3 Total Containers

C. Caution:

- 1. Extreme care must be taken to ensure that frozen samples do not break during shipment.
- Before adding soil to pre-weighed vials containing a stir bar, weigh the vials to confirm the tared weight. If the weight varies by more than 0.1 g, record the new weight on the label and the sample documentation. Do NOT add labels to these vials once the tared weight has been determined/confirmed.

D. Dry Samples:

All options include taking a sample in a dry 40 mL VOA vial (or a 4 oz wide mouth jar) with no headspace. No additional water, NaHSO₄, or methanol is added to this sample. This sample is taken to determine moisture content; therefore, it does not need to be tared or have a stir bar.

E. Iced or Frozen Samples:

- 1. Iced means cooled to 4° C ($\pm 2^{\circ}$ C) immediately after collection.
- 2. Frozen means cooled to between -7°C and -15°C immediately after collection.

F. Sample Delivery:

CLP strongly recommends that all samples reach the laboratory by COB the next day after sample collection.

G. Notes:

- 1. For Option 4, samples can be preserved with NaHSO₄ either:
 - In the field; or
 - In the laboratory upon receipt. In this case, the sampler should put the following information in the Preservation Column of the TR/COC Record "To be preserved at lab with NaHSO₄". This Regional Request should also be communicated to SMO so that the laboratory can be notified.
- 2. Regional QAPPs may require the use of Option 5. Please note that this option is for medium-level analysis ONLY.
- 3. If water, methanol, or NaHSO₄ preservative is added to the vials in the field, a field blank containing the appropriate liquid used in the vials should be sent to the laboratory for analysis.

H. Number of Containers Rationale:

The rationale for the number of containers (vials or samplers) required for the field sample and the required laboratory QC for each option is given as follows:

Option 1.

Rationale for Regular1 vial for low-level analysis (water purge)
1 vial for backup low-level analysis

1 vial for medium-level analysis (methanol extraction)

Rationale for QC2 vials for MS and MSD low-level analysis **Vials:**2 vials for MS and MSD medium-level analysis

2 vials for backup (MS and MSD) low-level or medium-level analysis

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Option 2.

Rationale for Regular 1 vial for low-level analysis (water purge) Vials:

1 vial for back up low-level analysis

1 vial dry for medium-level analysis (methanol extraction)

Rationale for QC 2 vials for MS and MSD low-level analysis Vials: 2 vials for MS and MSD medium-level analysis

2 vials for backup (MS and MSD) low-level or medium-level analysis

Medium-level: Methanol will be added in the laboratory

Analysis

Option 3.

Rationale for Regular 1 sampler for low-level analysis (water purge) 1 sampler for back up low-level analysis **Samples:**

1 sampler for medium-level analysis (methanol extraction)

Rationale for QC 2 samplers for MS and MSD low-level analysis

Samples: 2 samplers for backup MS and MSD low-level analysis

2 samplers for MS and MSD medium-level analysis

2 samplers for backup MS and MSD medium-level analysis

Option 4a (NaHSO₄ added in the field).

Rationale for Regular 1 vial with water for low-level analysis (water purge) Vials:

1 vial with water for backup low-level analysis

1 vial dry for medium-level analysis (methanol extraction)

Rationale for QC 2 vials with water for MS and MSD low-level analysis Vials: 2 vials dry for MS and MSD medium-level analysis

2 vials for backup (MS and MSD) low-level or medium-level analysis

Option 4b (NaHSO₄ added in the laboratory).

Rationale for Regular 1 vial for low-level analysis (water purge)

Vials: 1 vial for backup low-level analysis

1 vial for medium-level analysis (methanol extraction)

Rationale for OC 2 vials for MS and MSD low-level analysis Vials: 2 vials for MS and MSD medium-level analysis

2 vials for backup (MS and MSD) low-level or medium-level analysis

Option 5.

Rationale for Regular 1 vial for regular medium-level analysis

Samples: 1 vial for back up medium-level analysis

Rationale for QC 2 samples for MS and MSD

Samples: 2 samples for backup MS and MSD

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Option 6.

In this option, all Regular and QC samples for both low-level and medium analysis are taken as subsamples from the same container.

Rationale for Regular 1 glass jar for low-level analysis and medium-level analysis

Analysis 1 glass jar for backup low-level analysis and medium-level analysis

Rationale for 1 glass jar for low-level analysis and medium-level analysis

QC Analysis: 1 glass jar for backup low-level analysis and medium-level analysis

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Appendix C: General CLP Sample Collection Guidelines VOAs in Water



Regional guidance and/or specific Project Plan requirements will supersede the guidelines listed below.

Collect the following:

- At least two 40 mL glass containers with polytetrafluoroethylene (PTFE)-lined septa and open top screw-caps that are filled to capacity with no air bubbles, preserved to a pH of 2 with HCl, and cooled to 4°C (±2°C) immediately after collection. DO NOT FREEZE THE SAMPLES.
- If Selected Ion Monitoring (SIM) analysis is requested, at least two additional 40 mL glass containers with PTFE-lined septa and open top screw-caps that are filled to capacity with no air bubbles, preserved to a pH of 2 with HCl, and cooled to 4°C (±2°C) immediately after collection.

Test for Carbonates, Residual Chlorine, Oxidants, and Sulfides:

- It is very important that samplers obtain Regional guidance when testing and ameliorating for:
 - Carbonates:
 - Residual chlorine (e.g., municipal waters or industrial waste waters that are treated with chlorine prior to use or discharge); or
 - Oxidants.
- VOA samples containing carbonates react with the acid preservative causing effervescence (due to formation of carbon dioxide), which can cause loss of volatile analytes.
- Residual chlorine present in VOA samples can continue to react with dissolved organic matter. This continuous reaction may lead to inaccurate quantitation of certain analytes present in the sample at the time of collection.
- Residual chlorine and oxidants present in VOA samples can cause degradation of certain volatile analytes (e.g., styrene).

Perform the following for Pre-Preserved Vials:

- 1. Pour the sample slowly down the edge of the sample vial to avoid excess aeration or agitation of the sample during filling.
- 2. Fill the vial completely so that a reverse (convex) meniscus is present and ensure that there are no air bubbles present (either in the body or especially at the top of the vial).
- 3. Place the septum on the vial so that the PTFE side is in contact with the sample, and then firmly tighten the cap.
- 4. Gently flip the vial a few times to ensure that the sample is mixed with the acid preservative.
- 5. While holding the vial upright, gently tap the sample to check for air bubbles (either in the body or especially at the top of the vial).
- 6. If air bubbles are present, discard the sample and select a new vial in which to recollect a new sample. Repeat Steps 1 5 above.
- 7. Do NOT mix or composite samples for VOAs.
- 8. Cool sample to a temperature of 4°C (±2°C). Samplers should begin the cooling process in the field as samples are being collected. Double-bagged ice should be used. DO NOT FREEZE WATER SAMPLES.
- 9. Immediately transfer the vial to the sample shuttle (device that contains a "set" of VOA vials) once it has been collected. Do NOT allow ice to touch the vials.

Perform the Following for *Empty* Vials:

1. Rinse the vial with sample water prior to actual sample collection and preservation.



Regions vary in their approach to pre-rinsing and/or re-using sample vials (e.g., some Regions do not recommend pre-rinsing and/or re-use of pre-cleaned containers using sample water). Be sure to follow Regional guidance.

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- 2. Add 1-2 mL of acid preservative to the vial. Check to ensure that the sample you are collecting requires a preservative (follow Regional guidance).
- 3. Pour the sample slowly down the edge of the sample vial to avoid excess aeration and agitation of the sample.
- 4. Fill the vial completely so that a reverse (convex) meniscus is present and ensure that there are no air bubbles present (either in the body or especially at the top of the vial).
- 5. Place the septum on the vial so that the PTFE side is in contact with the sample, and then firmly tighten the cap.
- 6. Gently flip the vial a few times to ensure that the sample is mixed with the acid preservative.
- 7. While holding the vial upright, gently tap the vial to check for air bubbles (either in the body or especially at the top of the vial).
- 8. If air bubbles are present, discard the sample and recollect a new sample using the same sample vial. Repeat Steps 1 7 above.
- 9. Check the recollected sample for air bubbles. If air bubbles are present, additional sample water may be added to the vial to eliminate air bubbles. If there are air bubbles after three consecutive attempts to eliminate air bubbles by the addition of sample water, the entire sample and sample vial should be discarded and a new sample collected.
- 10. Do NOT mix or composite samples for VOAs.
- 11. Cool sample to a temperature of 4°C (±2°C). Samplers should begin the cooling process in the field as samples are being collected. Double-bagged ice should be used. DO NOT FREEZE WATER SAMPLES.
- 12. Immediately transfer the vial to the sample shuttle (device which contains a "set" of VOA vials) once it has been collected. Do NOT allow ice to touch the vials.

Things to Remember:

- Samples must be shipped as soon as possible, preferably on the same day as sample collection to avoid exceeding sample holding times. If overnight transit is not possible, samples should be maintained at 2 4°C until they are shipped to the laboratory.
- If samples are not preserved (a requirement for certain analytes), the technical holding time is shortened to 7 days.

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Appendix D: Sampling Techniques and Considerations

During a sampling event, the sampler is expected to follow prescribed sampling techniques. The sampler should also be aware of any special sampling considerations, contaminant issues, and sample compositing and mixing methods that could affect their sampling efforts.



Regional guidance will take precedence over any of the techniques and considerations listed below.

D.1 General Sampling Techniques

Information regarding surface water, sediment, soil, and groundwater sampling can be found in many documents including, but not limited to, the following sources:

- Compendium of ERT Surface Water and Sediment Sampling Procedures, EPA/540/P-91/005;
- Compendium of ERT Soil Sampling and Surface Geophysics Procedures, EPA/540/P-91/006;
- Compendium of ERT Groundwater Sampling Procedures, EPA/540/P-91/007;
- Quality Assurance Sampling Plan for Environmental Response (QASPER) software, Version 4.1, ERT; and
- Requirements for the Preparation of Sampling and Analysis Plans; United States Army Corps of Engineers, February 1, 2001, EM 200-1-3.

When working with potentially hazardous materials, samplers should follow USEPA and OSHA requirements, specific health and safety procedures, and DOT requirements.

D.2 Special Sampling Considerations

Samplers should refer to Regionally-developed SOPs to obtain specific procedures for properly collecting and preserving samples in the field. For additional guidance regarding sampling for VOAs in soil and water, see Appendices B and C. Samplers should obtain Regional guidance when testing and ameliorating for:

- Carbonates in VOA soil and water;
- Residual chlorine in VOA soil and water, or cyanide water;
- Oxidants in VOA soil and water; or
- Sulfides in cyanide.

D.3 Contaminant Sampling

Certain compounds can be detected in the parts-per-billion (ppb) and/or parts-per-trillion (ppt) range. Extreme care MUST be taken to prevent cross-contamination of these samples. The following precautions should be taken when trace contaminants are a concern:

- Disposable gloves should be worn each time a different location is sampled.
- When collecting both surface water and sediments, surface water samples should be collected first. This reduces the chance of sediment dispersal into surface water, and the resulting loss of surface water sample integrity.
- Sampling should occur in a progression from the least to the most contaminated area, if this information is known to the sampling team.
- Samplers should use equipment constructed of PTFE, stainless steel, or glass that has been properly pre-cleaned for collection of samples for trace organic and/or inorganic analyses. Equipment constructed of plastic or polyvinyl chloride (PVC) should NOT be used to collect samples for trace organic compound analyses.
- Equipment constructed of stainless steel should NOT be used to collect samples for trace metals analysis.

D.4 Sample Compositing

Sample compositing is a site-specific activity that must be conducted according to the SAP. Compositing is typically used for large sites under investigation to improve the precision (i.e., lower the variance) of the estimated average contaminant concentrations. Samples for VOA analysis should NOT be composited to minimize loss of VOAs/analytes.

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Composite samples consist of a series of discrete grab samples that are mixed together to characterize the average composition of a given material. The discrete samples are usually of equal volume, but may be weighted to reflect an increased flow or volume. Regardless, all discrete samples must be collected in an identical manner and the number of grab samples forming a composite should be consistent. There are several compositing techniques that may be required such as:

- Flow-proportioned Collected proportional to the flow rate during the compositing period by either a time-varying/constant volume or a time-constant/varying volume method. This technique is usually associated with wastewater or storm water runoff sampling.
- Time Composed of a varying number of discrete samples collected at equal time intervals during the compositing period. This technique is typically used to sample wastewater and streams, and in some air sampling applications.
- Areal Collected from individual grab samples collected in an area or on a cross-sectional basis. Areal composites are comprised of equal volumes of grab samples where all grabs are collected in an identical manner. This technique is typically used for estimating average contaminant concentrations in soils or sediments. This technique is useful when contaminants are present in nugget form (i.e., TNT chunks, lead shot, etc.), thus exhibiting large differences in concentration over a small sample area.
- Vertical Collected from individual grab samples but taken from a vertical cross section. Vertical composites are comprised of equal volumes of grab samples where all grab samples are collected in an identical manner. Examples would include vertical profiles of a soil borehole or sediment columns.
- Volume Collected from discrete samples whose aliquot volumes are proportional to the volume of sampled
 material. Volume composites are usually associated with hazardous waste bulking operations where the
 sample represents combined or bulked waste.

When compositing solid samples (i.e., sediment, soil, or sludge) for analysis of compounds present in trace quantities, use a stainless steel or PTFE bowl and spatula.

D.5 Sample Mixing and Homogenizing

Mixing of the sample for the remaining parameters is necessary to create a representative sample media. It is extremely important that solid samples be mixed as thoroughly as possible to ensure that the sample is as representative as possible of the sample location. Please refer to the project-specific SAP regarding instructions on removal of any extraneous materials (e.g., leaves, sticks, rocks, etc.). The mixing technique will depend on the physical characteristics of the solid material (e.g., particle size, moisture content, etc.). The mixing container should be large enough to hold the sample volume and accommodate the procedures without spilling. Both the mixing container (generally a bowl or tray) and the mixing implement should be properly decontaminated before use. Samples should be homogenized according to procedures listed in the project-specific SAP.

Samples for VOA analysis should not be mixed to minimize loss of volatile analytes.

Table D-1 provides a short procedure for mixing a soil sample with a small particle size (less than 1/4 in) and filling sample containers in the field.

Table D-1. Mixing a Sample and Filling Sample Containers

Step	Action
1	Roll the contents of the compositing container to the middle of the container and mix.
2	Quarter the sample and move to the sides of the container.
3	Mix each quarter individually, then combine and mix OPPOSITE quarters, then roll to the middle of the container.
4	Mix the sample once more, and then quarter the sample again.
5	Mix each quarter individually, then combine and mix ADJACENT corners, then roll to the middle of the container. The goal is to achieve a consistent physical appearance before sample containers are filled.
6	Flatten piled material into an oblong shape.
7	Using a flat-bottomed scoop, collect a strip of soil across the entire width of the short axis and place it into a sample container.
8	Repeat Step 7 at evenly-spaced intervals until the sample containers are filled.
9	Record the approximate quantity of each subsample in the field log book.

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Appendix E: Sampling Checklists

Appendix E-1: Personnel Preparation Checklist (Page 1 of 1)

	Personnel Briefing			Comments:
1.	$\label{thm:polynomial} \mbox{Did you review sampling team responsibilities and identify individual(s) responsible for corrective actions?}$			
2.	Did you ensure that you have met the appropriate personal safety and protection requirements?			
3.	Did you identify sampling locations and receive permission to access them, as appropriate?			
4.	Did you contact the appropriate utility companies PRIOR to the start of sampling?			
No	By law, utility companies must be contacted prior to the start of digging/sampling so that any underground utilities (gas lines, water lines, electrical lines, etc.) can be marked. A list of one-call centers for each state may be found at: http://www.digsafely.com/contacts.htm .			
5.	If sampling on private property, do you have sample receipts to provide to the property owner for all samples taken and removed from the property?			
6.	Have you determined the number and type of samples to be collected?			
7.	Did you review sample collection methods?			
8.	Have you reviewed sample container requirements?			
9.	Did you review decontamination requirements, procedures, and locations?			
10.	Did you determine holding times and conditions?			
11.	Did you determine Performance Evaluation (PE) and Quality Control (QC) sample requirements?			
12.	Have you obtained shipping cooler temperature blanks, if required?			
13.	Did you review sample label and tag requirements?			
14.	Did you review Traffic Report/Chain of Custody (TR/COC) Record and custody seal requirements?			
15.	Have you obtained the laboratory name, shipping addresses, and telephone number?			
16.	Did you review cooler return instructions?			
17.	Have you obtained shipping company information (name, telephone number, account number, pickup schedule)?			
18.	Have you obtained shipping schedules?			
19.	Did you review shipment reporting requirements and the appropriate contact names and telephone numbers for reporting?			
20.	Have you included any sampler comments regarding sampling issues (e.g., low volumes, matrix, suspected concentrations based on field measurements)?			

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Appendix E-2: General Sample Collection Checklist $(Page\ 1\ of\ 1)$

	General Sample Collection	Yes	No	Comments:
1.	Did you identify and mark the sampling location with buoys, flags, or stakes according to the sampling plans, maps, and grids?			
2.	If the sampling location is inaccessible, did you contact the appropriate field or Regional personnel for instructions?			
3.	Did you use the correct sampling equipment?			
4.	Did you follow the correct decontamination procedures?			
5.	Did you follow the correct collection procedures?			
6.	Did you use the correct sample containers for each sample collected?			
7.	Did you collect the correct volume for each sample?			
8.	8. Did you collect the correct type of sample, including primary samples and Quality Control (QC) samples?			
9.	Did you properly preserve each sample collected?			
10.	Did you correctly document and label each sample with all necessary information?			
	Under no circumstances should the site name appear on any documentation being sent to the laboratory.			
11.	If sampling on private property, did you provide a sample receipt to the owner of the property for all samples taken and removed from the property?			

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Appendix E-3: Completing Field Logbook Checklist (Page 1 of 1)

	Completing Field Logbook	Yes	No	Comments:
1.	Did you use waterproof ink when writing in the field logbook?			
2.	 Did you document sampling project information such as: Project name, ID, and location; Names of samplers; Geological observations, including maps; Atmospheric conditions; Field measurements; and Sampling dates, times, and locations? Under no circumstances should the site name appear on any documentation being sent to the laboratory.			
3.	Did you record sampling activity information such as: Sampling dates and times; Sample identifications; Sample matrices; Sample descriptions (e.g., odors and/or colors); Number of samples taken; Sampling methods/equipment; and Description of QC samples?			
4.	Did you document any and all deviations from the sampling plan?			
5.	Did you document any and all difficulties in sampling and/or any unusual circumstances?			
6.	Were all errors corrected by crossing a line through the error, initialing the error, dating the error, and then adding the correct information?			

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Appendix E-4: Completing Handwritten Sample Labels Checklist $(Page\ 1\ of\ 1)$

	Completing Handwritten Sample Labels	Yes	No	Comments:
1.	Did the Region provide CLP Sample Numbers and SMO-assigned Case Numbers?			
2.	If additional CLP Sample Numbers were needed, did you contact the appropriate Regional personnel?			
3.	Were the CLP Sample Numbers and SMO-assigned Case Numbers on the labels correct? Organic CLP Sample Numbers begin with the Regional letter code, followed by letters and numbers. Inorganic CLP Sample Numbers begin with "M", followed by the Regional letter code, and then letters and numbers.			
	The following characters are not used in generating CLP Sample Numbers and should never appear on any paperwork send to the laboratory: I; O; U; and V. Also, the last character of a CLP Sample Number will never be a letter.			
4.	Were samples uniquely numbered and designated to only one sample?			
	Samples collected for total metal and dissolved metal analyses must receive separate, unique, CLP Sample Numbers.			
5.	Were Quality Control (QC) samples numbered accordingly?			
6.	Were the specific requirements followed for total and dissolved metals analysis, QC and Performance Evaluation (PE) samples, and SW-846 Method 5035A?			
7.	Were all temperature blanks labeled with "TEMPERATURE BLANK"?			
8.	Was a sample label containing the CLP Sample Number, SMO-assigned Case Number, location, concentration, preservative, and the fraction/analysis, attached to each sample bottle or container as the sample was collected?			
	Under no circumstances should the site name appear on any documentation being sent to the laboratory.			
9.	Was clear tape placed over the sample labels to protect the labels from moisture and to help the labels adhere to the sample bottle?			
10.	Were all errors corrected by crossing a line through the error, initialing the error, dating the error, and then adding the correct information?			

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Appendix E-5: Completing Handwritten Sample Tags & Custody Seals Checklists $(Page\ 1\ of\ 1)$

	Completing Handwritten Sample Tags	Yes	No	Comments:
1.	Was waterproof ink used on the sample tags?			
2.	If Regionally required for individual sample containers, was the project code on the sample tag completed?			
3.	Was the station number on the sample tag completed?			
4	Was the date filled in using the format MM/DD/YYYY?			
5.	Was the time of sample collection indicated in military time format HH:MM?			
6.	Was the box checked indicating composite or grab sample?			
7.	Was the station location on the sample tag completed?			
8.	Did you indicate whether or not the sample was preserved by checking "yes" or "no?"			
9.	Was the appropriate analysis indicated on the sample tag?			
10.	Were the appropriate CLP Sample Number and SMO-assigned Case Number indicated and cross-referenced with the numbers on the sample label?			
11.	Did you sign the sample tags?			
12	Did you attach the sample tag to the neck of the sample bottle with striang, stretch string, or wire (recommended method)?			
(Do NOT use wire to attach a sample tag to a metal sample.			
13.	Were all errors corrected by crossing a line through the error, initialing the error, dating the error, and then adding the correct information?			
	Completing Custody Seals	Yes	No	Comments:
1.	Did you sign and date the custody seal?			
2.	Did you attach a completed custody seal to the sample bottle, container, or plastic bag, placing the seal over the cap or lid of each sample bottle or container or on the bag opening such that it will be broken if the sample bottle, container, or bag is opened or tampered with?			
3.	As appropriate, did you attach the completed custody seal to the sample shipping container or cooler, placing the seal such that it will be broken if the container or cooler is opened or tampered with?			
4.	Were all errors corrected by crossing a line through the error, initialing the error, dating the error, and then adding the correct information?			

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Appendix E-6: Packing Sample Container Checklist (Page 1 of 1)

	Packing Sample Container	Yes	No	Comments:
	1. Did you follow all State, Federal, Department of Transportation (DOT), and International Air Transportation Association (IATA) regulations governing the packaging of environmental and hazardous samples?			
No	If samples contain methanol preservation (e.g., samples to be analyzed by SW-846 Method 5035A), refer to the packaging instructions in Appendix A.			
2.	Were all CLP Sample Numbers, SMO-assigned Case Numbers, fractions/analyses, labels, tags, and custody seals attached to the correct sample containers?			
3.	Was an inventory conducted of CLP Sample Numbers, SMO-assigned Case Numbers, fractions/analyses, and containers, and verified against the TR/COC Records?			
4.	Were the correct number and type of Performance Evaluation (PE) and Quality Control (QC) samples collected?			
5.	Were all sample containers sealed in clear plastic bags with the sample label and tag visible through the packaging?			
6.	Were all soil/sediment samples known to contain dioxin securely enclosed in metal cans (e.g., paint cans) with the lids sealed?			
7.	Was suitable absorbent packing material placed around the sample bottles or containers?			
8.	Were the outsides of metal containers labeled properly with the CLP Sample Number, SMO-assigned Case Number, and the fraction/analysis of the sample inside?			

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Appendix E-7: Packing Shipping Container Checklist (Page 1 of 1)

	Packing Shipping Container	Yes	No	Comments:
1.	Were you shipping samples in a clean waterproof metal or hard plastic ice chest or cooler in good condition?			
2.	Were all non-applicable labels from previous shipments removed from the container?			
3.	Were all inside and outside drain plugs closed and covered with suitable tape (e.g., duct tape)?			
4.	Was the inside of the cooler lined with plastic (e.g., large heavy-duty garbage bag)?			
5.	Was the lined shipping cooler packed with noncombustible absorbent packing material?			
6.	Were sample containers placed in the cooler in an upright position not touching one another?			
7.	Was a sample shipping cooler temperature blank included in the cooler?			
8.	Did the documentation in the cooler only address the samples in that cooler?			
9.	Was the site name absent from all documentation?			
	Under no circumstances should the site name appear on any documentation being sent to the laboratory.			
10.	Was there sufficient packing material around and in between the sample bottles and cans to avoid breakage during transport?			
11.	If required, was double-bagged ice placed on top and around sample bottles to keep the samples cold at 4° C (\pm 2° C)?			
	Do Not Pack Loose Ice Into the Cooler!			
12.	Was the top of the plastic liner fastened and secured with tape?			
13.	Was a completed custody seal placed around the top of the fastened plastic liner (if required by the Region)?			
14.	Were all sample documents enclosed within the cooler (e.g., TR/COC Record and cooler return instructions) in a waterproof plastic bag?			
15.	Was the plastic bag, containing the documentation, taped to the underside of the cooler lid?			
16.	Were cooler return instructions and airbills, if required, taped to the underside of the cooler lid?			
17.	Was the return address of the cooler written with permanent ink on the underside of the cooler lid?			
18.	Was tape placed around the outside of the entire cooler and over the hinges?			
19.	Were the completed custody seals placed over the top edge of the cooler so the cooler cannot be opened without breaking the seals?			
20.	Was the return address label attached to the top left corner of the cooler lid?			
21.	Were instructional labels attached to the top of the cooler, as necessary (e.g., "This End Up," "Do Not Tamper With," or "Environmental Laboratory Samples")?			
22.	If shipping hazardous samples, were the correct labels attached to the cooler (e.g., "Flammable Liquids", "Caution", or "Poison")?			
23.	If shipping samples containing methanol as a preservative (e.g., samples to be analyzed by SW-846 Method 5035A), was a label used to indicate methanol, the United Nations (UN) identification number for methanol (UN 1230), and Limited Quantity?			

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Appendix E-8: Shipping & Reporting CLP Samples Checklist $(Page\ 1\ of\ 1)$

	Shipping CLP Samples	Yes	No	Comments:
1.	Did you follow all State, Federal, Department of Transportation (DOT), and International Air Transportation Association (IATA) regulations governing the shipment of environmental and hazardous samples?			
2.	Was a separate airbill filled out for each cooler being shipped?			
3.	Was the airbill filled out completely, including correct laboratory name, address, and telephone number, identification of recipient as "Sample Custodian," and appropriate delivery option (e.g., overnight or Saturday)?			
4.	Was the completed airbill attached to the top of the cooler with the correct laboratory address?			
5.	If more than one cooler was being shipped to the same laboratory, were they marked as "1 of 2," "2 of 2," etc.?			
6.	Were the samples being shipped "overnight" through a qualified commercial carrier?			
	Reporting CLP Samples	Yes	No	Comments:
1.	Did you contact the Contract Laboratory Program Sample Management Office (SMO) on the same day samples were shipped?			
2.	If the samples were shipped after 5:00 PM Eastern Time (ET), were they reported to the RSCC (or designee) or to SMO by 8:00 AM ET the following business day?			
3.	Did you notify the RSCC (or designee) or SMO so that SMO will receive the delivery information by 3:00 PM ET on Friday for sample shipments that will be delivered to the laboratory on Saturday?			
4.	 Did you provide the RSCC (or designee) or SMO with: Your name, phone number, and Region number; Case Number of the project; Exact number of samples, matrix(ces), concentration(s), and type of analysis; Laboratory(ies) to which the samples were shipped; Carrier name and airbill number; Date of shipment; Date of next shipment; and Any other information pertinent to the shipment? 			

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Appendix F: Glossary

Analyte -- The element, compound, or ion that is determined in an analytical procedure; the substance or chemical constituent of interest.

Analytical Services Branch (**ASB**) -- Directs the Contract Laboratory Program (CLP) from within the United States Environmental Protection Agency's (USEPA's) Office of Superfund Remediation and Technology Innovation (OSRTI) in the Office of Solid Waste and Emergency Response (OSWER).

Aroclor -- Polychlorinated biphenyls (PCBs) or a class of organic compounds with 1 to 10 chlorine atoms attached to biphenyl and a general chemical formula of $C_{12}H_{10-x}Cl_x$. PCBs, commercially produced as complex mixtures containing multiple isomers at different degrees of chlorination, were marketed in North America under the trade name Aroclor.

Case -- A finite, usually predetermined, number of samples collected over a given time period from a particular site. Case Numbers are assigned by the Sample Management Office (SMO). A Case consists of one or more Sample Delivery Groups (SDGs).

Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) -- Initiated in December 1980, CERCLA provided broad federal authority to respond directly to the release or possible release or hazardous substances that may endanger human health or the environment. CERCLA also established a trust fund to provide for cleanup when no responsible party could be identified; hence CERCLA is commonly referred to as "Superfund".

Contract Laboratory Program (CLP) -- A national program of commercial laboratories under contract to support the USEPA's nationwide efforts to clean up designated hazardous waste sites by providing a range of chemical analytical services to produce environmental data of known and documented quality. This program is directed by USEPA's Analytical Services Branch (ASB).

Contract Laboratory Program Project Officer (CLP PO) -- Monitors technical performance of the contract laboratories in each Region.

Contract Laboratory Program Sample Management Office (CLP SMO) -- A contractor-operated facility operated under the CLP, awarded and administered by the USEPA, which provides necessary management, operations, and administrative support to the CLP. SMO coordinates and schedules sample analyses, tracks sample shipments and analyses, receives and tracks data for completeness and compliance, and processes laboratory invoices.

Custody Seal -- An adhesive label or tape that is used to seal a sample bottle or container that maintains chain-of-custody and that will break if the sample bottle or container is opened or tampered with.

Cyanide (**Total**) -- Cyanide ion and complex cyanides converted to hydrocyanic acid (HCN) by reaction in a reflux system of a mineral acid in the presence of magnesium ion.

Data Quality Objective (DQO) -- The requirements established to maintain the quality of the data being collected.

Data Validation -- Data validation is based on Region-defined criteria and limits, professional judgment of the data validator, and (if available) the Quality Assurance Project Plan (QAPP) and Sampling and Analysis Plan (SAP).

Equipment Blank -- A sample used to check field decontamination procedures. See Field Blank.

Field Blank -- Any blank sample that is submitted from the field. Each field blank is assigned its own unique USEPA Sample Number. A Field Blank checks for cross-contamination during sample collection, sample shipment, and in the laboratory. A field blank includes trip blanks, rinsates, equipment blanks, etc.

Field Duplicate -- Checks reproducibility of laboratory and field procedures and indicates non-homogeneity.

Field Operations Reporting Management System (FORMS) II Lite -- A stand-alone, Windows-based software application that enables samplers to automatically create and generate sample documentation both prior to and during a sampling event.

Field QC Sample -- Used to detect for contamination or error in the field.

Field Sample -- Primary sample material taken out in the field from which other samples, such as duplicates or split samples are derived. A field sample can be prepared in the field and sent for analysis in one or multiple containers, and is identified by a unique EPA Sample Number.

Field Sampling Plan (FSP) -- Developed to outline the actual steps and requirements pertaining to a particular sampling event, and explains, in detail, each component of the event to all involved samplers.

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Holding Time -- The elapsed time expressed in hours, days, or months from the date of collection of the sample until the date of its analysis.

Contractual -- The lengths of time that the CLP laboratory must follow to comply with the terms of the contract, and are described in the CLP analytical services Statements of Work (SOWs).

Technical -- The maximum lengths of time that samples may be held from time of collection to time of preparation and/or analysis and still be considered valid.

Laboratory Blank -- See Method Blank.

Laboratory Duplicate -- A sample required by the laboratory's contract to check the precision of inorganic analyses.

Laboratory QC Sample -- An additional volume of an existing sample, as required by the laboratory's contract, used to detect contamination or error in the laboratory's practices.

Matrix -- The predominant material of which a sample to be analyzed is composed.

Matrix Spike (MS) -- Sample required by the laboratory's contract to check the accuracy of organic and inorganic analyses. It is an aliquot of a sample (water or soil) that is fortified (spiked) with known quantities of a specific compound and subjected to the entire analytical procedure. See Matrix Spike Duplicate.

Matrix Spike Duplicate (MSD) -- Sample required by the laboratory's contract to check the accuracy and precision of organic analyses. It is a second aliquot of the same matrix as the Matrix Spike (MS) that is spiked to determine the precision of the method. See Matrix Spike.

Method Blank -- An analytical control consisting of all reagents, internal standards and surrogate standards [or System Monitoring Compounds (SMCs) for volatile organic analysis], that is carried throughout the entire analytical procedure. The method blank is used to define the level of laboratory, background, and reagent contamination, also referred to as laboratory blank when defining the level of laboratory contamination.

Performance Evaluation (PE) Sample -- A sample of known composition provided by the USEPA for contractor analysis. Used by USEPA to evaluate contractor performance.

Pesticides -- Substances intended to repel, kill, or control any species designated a "pest", including weeds, insects, rodents, fungi, bacteria, and other organisms. Under the CLP, only organochlorine pesticides are analyzed (e.g., DDT, Dieldrin, Endrin, etc.).

Polychlorinated Biphenyls (PCBs) -- A group of toxic, persistent chemicals used in electrical transformers and capacitors for insulating purposes, and in gas pipeline systems as a lubricant. The sale and new use of PCBs were banned by law in 1979.

Quality Assurance (QA) -- An integrated system of management activities involving planning, implementation, assessment, reporting, and quality improvement to ensure that a process, item, or service is of the type and quality needed and expected by the customer.

Quality Assurance Project Plan (QAPP) -- Document written to meet requirements outlined in the document *EPA Guidance for Quality Assurance Project Plans* (EPA QA/R-5). Prepared in advance of field activities and used by samplers to develop any subsequent plans such as the Sampling Analysis Plan (SAP) or the Field Sampling Plan (FSP).

Quality Control (QC) -- The overall system of technical activities that measures the attributes and performance of a process, item, or service against defined standards to verify that they meet the stated requirements established by the customer; operational techniques and activities that are used to fulfill requirements for quality.

Regional Sample Control Center (RSCC) Coordinator -- In most Regions, coordinates sampling efforts and serves as the central point-of-contact for sampling questions and problems. Also assists in coordinating the level of Regional sampling activities to correspond with the monthly projected demand for analytical services.

Regional Site Manager -- Coordinates the development of data quality objectives and oversees project-specific remedial or removal contractors, State officials, or private parties conducting site sampling efforts.

Rinse Blank -- A sample used to check decontamination procedures. Also see Field Blank.

Routine Analytical Service (RAS) -- The standard inorganic and organic analyses available through the CLP.

Sample -- A discrete portion of material to be analyzed that is contained in single or multiple containers, and identified by a unique Sample Number.

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Sample Delivery Group (SDG) – A unit within a sample Case that is used to identify a group of samples for delivery. An SDG is defined by the following, whichever is most frequent:

- Each Case of field samples received; or
- Each 20 field samples (excluding PE samples) within a Case; or
- Each 7 calendar day period (3 calendar day period for 7-day turnaround) during which field samples in a Case are received (said period beginning with the receipt of the first sample in the SDG).

In addition, all samples and/or sample fractions assigned to an SDG must have been scheduled under the same contractual turnaround time. Preliminary Results have no impact on defining the SDG. Sample may be assigned to SDGs by matrix (e.g., all soil samples in one SDG, all water samples in another) at the discretion of the laboratory.

Sample Label -- An identification label attached to a sample bottle or container to identify the sample.

Sample Number -- A unique number used to identify and track a sample. This number can be recorded on a sample label or written on the sample bottle or container using indelible ink.

Sample Tag -- A tag attached to a sample that identifies the sample and maintains chain-of-custody.

Sampling Analysis Plan (SAP) -- A document that explains how samples are to be collected and analyzed for a particular sampling event.

Semivolatile Organic Analyte (SVOA) -- A compound amenable to analysis by extraction of the sample using an organic solvent.

Statement of Work (SOW) -- A document that specifies how laboratories analyze samples under a particular Contract Laboratory Program (CLP) analytical program.

Superfund -- The program operated under the legislative authority of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) and Superfund Amendments and Reauthorization Act (SARA) that funds and carries out USEPA removal and remedial activities at hazardous waste sites. These activities include establishing the National Priorities List (NPL), investigating sites for inclusion on the list, determining their priority, and conducting and/or supervising cleanup and other remedial actions.

Superfund Amendments and Reauthorization Act (SARA) -- The 1986 amendment to the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA).

Traffic Report/Chain of Custody (TR/COC) Record -- A record that is functionally similar to a packing slip that accompanies a shipment of goods. Used as physical evidence of sample custody and functions as a permanent record for each sample collected.

Trip Blank -- A sample used to check for contamination during sample handling and shipment from field to laboratory. Also see Field Blank.

Volatile Organic Analyte (VOA) -- A compound amenable to analysis by the purge-and-trap technique. Used synonymously with the term purgeable compound.

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APPENDIX B SITE-SPECIFIC DATA QUALITY OBJECTIVES

STEP 1. STATE THE PROBLEM

Uranium and other metals/radionuclides from mine tailings may be migrating to surface water and/or groundwater at the former Jackpile Mine representing a threat to human health and the environment.

STEP 2. IDENTIFY THE DECISION

Groundwater samples will be collected from upgradient, on-site, and downgradient of the mine area to determine whether contaminants of concern (CoCs) identified during the 2010 SI sampling event are significantly elevated in groundwater with respect to background concentrations. In addition, general water chemistry analyses are needed to update a hydrologic model of the site.

update a hydrologic model of the site.	
IDENTIFY THE ALTERNATIVE ACTIONS THAT MAY BE TAKEN BASED ON THE DECISIONS.	 If concentrations of CoCs are significantly elevated with respect to background in groundwater monitoring wells constructed on-site, and downgradient of, the site, then an observed release to groundwater has been established. If concentrations of CoCs in groundwater samples are not significantly elevated with respect to background,
	then an area of observed contamination has not been established.
	General water chemistry data will be used to update the hydrologic model of the site.
STEP 3. IDENTIFY INPUTS TO THE DECISION	
IDENTIFY THE INFORMATIONAL INPUTS NEEDED TO RESOLVE A DECISION.	 Review historical site operations contributing to contamination. Historical groundwater and surface water data collected at the site.
	 Contaminant concentrations in waste soil samples collected during 2010 SI sampling. New groundwater sampling during this ESI.
IDENTIFY THE SOURCES FOR EACH INFORMATIONAL INPUT AND LIST THE INPUTS	Existing groundwater monitoring well locations as shown in Figure 3-1.
THAT ARE OBTAINED THROUGH ENVIRONMENTAL MEASUREMENTS.	 Inorganic chemical analyses for Total Dissolved Solids by EPA Method 160.1, TAL metals by 6010/7470, and General Anion Chemistry by EPA Method 9056. Radionuclide analysis for uranium isotopic species by Alpha Spec ASTM 3972-90M, Gross Alpha, Beta, and
	Gamma by EPA Method 9310, and Radium 226 +228 by EPA Method 903.1.
BASIS FOR THE CONTAMINANT SPECIFIC ACTION LEVELS.	For an observed release, contaminants must be detected at concentrations greater than three-times (>3X) background for an analyte that is detected in background samples, or detected where the analyte is not detected in background samples.

IDENTIFY POTENTIAL SAMPLING TECHNIQUES AND APPROPRIATE ANALYTICAL METHODS.	 Groundwater samples will be collected in accordance with US EPA Groundwater Sampling Compendium, OSWER 9360.4-06 and Weston Groundwater Sampling SOP 1002-02. Inorganic chemical analyses for Total Dissolved Solids by EPA Method 160.1, TAL metals by EPA Method 6010/7470, and General Cation/Anion Chemistry by EPA Method 9056. Radionuclide analysis for uranium isotopic species by Alpha Spec ASTM 3972-90M, Gross Alpha, Beta, and Gamma by EPA Method 9310, and Radium 226 + 228 by EPA Method 903.1.
STEP 4. DEFINE THE BOUNDARIES OF THE STU	JDY
DEFINE THE DOMAIN OR GEOGRAPHIC AREA WITHIN WHICH ALL DECISIONS MUST APPLY.	Selected groundwater monitoring wells located on site, up to one mile up gradient, and up to five miles downgradient of the site.
SPECIFY THE CHARACTERISTICS THAT DEFINE THE POPULATION OF INTEREST.	 Groundwater includes the Alluvial Aquifer, which is associated with the active surface water system, and the Jackpile Aquifer, which is partially confined and has a distinct water chemistry.
DEFINE THE SCALE OF DECISION MAKING.	Results of groundwater and surface water sampling will be used to evaluate the groundwater-to-surface water pathway to the HRS evaluation.
DETERMINE THE TIME FRAME TO WHICH THE DATA APPLY.	The data will apply until the site media, represented by the waste, groundwater, and surface water samples, receives appropriate remedial actions.
DETERMINE WHEN TO COLLECT DATA.	Samples will be collected during the START-3 field effort and prior to HRS evaluation. This should take place during the period of low evapotranspiration – during the months of October to April.
IDENTIFY PRACTICAL CONSTRAINTS ON DATA COLLECTION.	 Inclement weather. Access not attainable. Wells destroyed or inoperable Vicious attack elk
STEP 5. DEVELOP A DECISION RULE	
SPECIFY THE PARAMETER THAT CHARACTERIZES THE POPULATION OF INTEREST.	Detection of CoCs in the water samples by analytical testing to confirm concentrations in the pathway that are significantly elevated with respect to background.

SPECIFY THE ACTION LEVEL FOR THE DECISION.	 The results of the 2010 SI indicate that COCs identified in the waste soil samples are CERCLA hazardous substances as defined in CERCLA Sections 101(14) and 101(33); therefore, an area of observed contamination has been established. If COCs are detected in on-site and downgradient groundwater at concentrations that are significantly elevated with respect to background, then an observed release has been established. The data will contribute to the HRS scoring. If COCs in the groundwater samples are either not detected at concentrations significantly elevated with respect to background, then an observed release has not been established. The surface water and/or groundwater-to-surface water pathways will not contribute to the HRS scoring.
STEP 6. SPECIFY LIMITS ON DECISION ERRORS	S
DEVELOP A DECISION RULE.	If site and/or downgradient groundwater samples have CoCs that are detected at elevations that are significantly above background concentrations, then an observed release to the groundwater pathway has been established.
DETERMINE THE POSSIBLE RANGE OF THE PARAMETER OF INTEREST.	Contaminant concentrations may range from 0 µg/mg to more than the contaminant specific action level.
DEFINE BOTH TYPES OF DECISION ERRORS AND IDENTIFY THE POTENTIAL CONSEQUENCES OF EACH.	Type I Error: Deciding that the specified area represented by the groundwater sample does not exceed the specified assessment level when, in truth, the concentration of the contaminant exceeds its specified assessment level. The consequence of this decision error is that remedial efforts in the watershed may not be undertaken, possibly endangering human health and the environment. This decision error is more severe.
	Type II Error: Deciding that the specified area represented by the groundwater sample does exceed the specified assessment level when, in truth, it does not. The consequences of this decision error are that remediation of the watershed will continue and unnecessary costs will be incurred.
ESTABLISH THE TRUE STATE OF NATURE FOR EACH DECISION RULE.	The true state of nature when the groundwater is decided to be below the specified assessment levels when in fact, it is not below the specified assessment levels, is that the watershed does need remedial action. The true state of nature when the groundwater is decided to be above the specified assessment levels when in fact, it is not above the specified assessment levels, is that the watershed does not need remedial action.

STEP 6. SPECIFY LIMITS ON DECISION ERRORS (Continued)					
DEFINE THE TRUE STATE OF NATURE FOR THE MORE SEVERE DECISION ERROR AS THE BASELINE CONDITION OR THE NULL HYPOTHESIS (H_o) AND DEFINE THE TRUE STATE FOR THE LESS SEVERE DECISION ERROR AS THE ALTERNATIVE HYPOTHESIS (H_a) .	Ho: The water represented by the sample is above the specified action level. Ha: The water represented by the sample is below the specified action level.				
ASSIGN THE TERMS "FALSE POSITIVE" AND "FALSE NEGATIVE" TO THE PROPER DECISION ERRORS.	 False Positive Error = Type I False Negative Error = Type II 				
ASSIGN PROBABILITY VALUES TO POINTS ABOVE AND BELOW THE ACTION LEVEL THAT REFLECT THE ACCEPTABLE PROBABILITY FOR THE OCCURRENCES OF DECISION ERRORS.	To be assigned based on discussions with EPA SAM.				
STEP 7. OPTIMIZE THE DESIGN					
REVIEW THE DQOs.	Due to insufficient historical data, determination of the standard deviation was not possible. Therefore, sample size calculation using the traditional statistical formula may not be the optimal design. In order to select the optimal sampling program that satisfies the DQOs and is the most resource effective, other elements were considered.				

DEVELOP GENERAL SAMPLING AND ANALYSIS DESIGN.

Up to fourteen groundwater samples will be collected from existing monitoring wells completed at locations upgradient, on-site, and downgradient of the Jackpile-Paguate Uranium Mine site. The samples will be analyzed for CoCs to determine the presence of hazardous substances as defined by CERCLA, and to show potential attribution to the pathways of concern used for HRS evaluation. The samples will be analyzed for uranium isotopes, TAL metals, gross alpha, beta, and gamma, and radium. In addition, water samples will be analyzed for general cation/anion chemistry and total dissolved solids.

DATA QUALITY OBJECTIVE NO. 2 JACKPILE-PAGUATE URANIUM MINE MEDIA OF CONCERN: SURFACE WATER/SEDIMENT

STEP 1. STATE THE PROBLEM

Are surface water/sediment samples collected from the surface water pathway above three times the maximum background concentrations resulting in an observed release?

STEP 2. IDENTIFY THE DECISION

Are the concentrations of chemicals of concern in surface water/sediment, represented by a sample, above three times the maximum background concentrations?

IDENTIFY THE ALTERNATIVE ACTIONS THAT MAY BE TAKEN BASED ON THE DECISIONS.

- If any contaminant exceeds three times the background concentrations in the surface water/sediment, the surface water/sediment represented by that sample will be considered contaminated and will require additional attention.
- If no contaminants exceed three times the background concentrations in the surface water/sediment, the surface water/sediment represented by that sample will not require additional attention.
- General water chemistry data from surface water will be used to update the hydrologic model of the site.

STEP 3. IDENTIFY INPUTS TO THE DECISION

IDENTIFY THE INFORMATIONAL INPUTS NEEDED TO RESOLVE A DECISION.

- Review historical site operations contributing to contamination.
- Historical surface water data collected at the site.
- Contaminant concentrations in waste soil samples and surface water samples collected during 2010 SI sampling.
- New groundwater sampling during this ESI.

IDENTIFY THE SOURCES FOR EACH INFORMATIONAL INPUT AND LIST THE INPUTS THAT ARE OBTAINED THROUGH ENVIRONMENTAL MEASUREMENTS.

- Surface water/Sediment samples from the surface water pathway.
- Inorganic chemical analyses for Total Dissolved Solids by EPA Method 160.1, and General Anion Chemistry by EPA Method 9056.
- Radionuclide analysis for Gross Alpha, Beta, and Gamma by EPA Method 9310, and Radium 226 +228 by EPA Method 903.1.

BASIS FOR THE CONTAMINANT SPECIFIC ACTION LEVELS.

The site specific screening levels will be three times the maximum background concentrations.

IDENTIFY POTENTIAL SAMPLING TECHNIQUES AND APPROPRIATE ANALYTICAL METHODS.

- Grab surface water/sediment samples from the surface water pathway analyzed for:
- General Anion Chemistry by EPA Method 9056
- Gross Alpha, Beta, and Gamma by EPA Method 9310
- Radium 226 +228 by EPA Method 903.1

STEP 4. DEFINE THE BOUNDARIES OF THE STUDY

DEFINE THE DOMAIN OR GEOGRAPHIC AREA WITHIN WHICH ALL DECISIONS MUST APPLY.

The boundaries for the Jackpile-Paguate Uranium Mine site are shown along with the proposed surface water/sediment sample locations in Figure 3-1.

DATA QUALITY OBJECTIVE NO. 2 JACKPILE-PAGUATE URANIUM MINE MEDIA OF CONCERN: SURFACE WATER/SEDIMENT

SPECIFY THE CHARACTERISTICS THAT DEFINE THE POPULATION OF INTEREST.	Contaminant concentrations in surface water/sediments at the sample locations.		
DEFINE THE SCALE OF DECISION MAKING.	The scale of decision will be for the site activities occurring at the time of the sample collection.		
DETERMINE THE TIME FRAME TO WHICH THE DATA APPLY.	The analytical data will apply until the surface water/sediment represented by the sample receives appropriate response action.		
STEP 4. DEFINE THE BOUNDARIES OF THE STU	JDY (Continued)		
DETERMINE WHEN TO COLLECT DATA.	Surface water/sediment samples will be collected during the field sampling activities.		
IDENTIFY PRACTICAL CONSTRAINTS ON DATA COLLECTION.	Inclement weather.Access not attainable.		
STEP 5. DEVELOP A DECISION RULE			
SPECIFY THE PARAMETER THAT CHARACTERIZES THE POPULATION OF INTEREST.	The sample concentrations at each sample location will be compared to three times the maximum background concentrations.		
SPECIFY THE ACTION LEVEL FOR THE DECISION.	The site specific screening levels will be three times the maximum background concentrations.		
DEVELOP A DECISION RULE.	If any result in a surface water/sediment sample is above three times the maximum background concentrations, then the surface water/sediment represented by that sample will require additional attention, otherwise the surface water/sediment does not require additional attention. Additional attention means more sampling, surface water/sediment collection and treatment, or other action deemed necessary by EPA.		
STEP 6. SPECIFY LIMITS ON DECISION ERROR	s		
DETERMINE THE POSSIBLE RANGE OF THE PARAMETER OF INTEREST.	Contaminant concentrations may range from 0 mg/kg for surface water/sediment to more than three times the maximum background concentrations.		
DEFINE BOTH TYPES OF DECISION ERRORS AND IDENTIFY THE POTENTIAL CONSEQUENCES OF EACH.	Type I Error: Deciding that the specified area represented by the surface water/sediment sample does not exceed three times the maximum background concentrations when, in truth, the surface water/sediment concentration of the contaminant exceeds three times the maximum background concentrations. The consequence of this decision error is that contaminated surface water/sediment will remain in place, possibly endangering human health and the environment. This decision error is more severe. Type II Error: Deciding that the specified area represented by the surface water/sediment sample does exceed three times the maximum background concentrations when, in truth, it does not. The consequences of this decision error		
	are that remediation of the surface water/sediment will continue and unnecessary costs will be incurred.		

DATA QUALITY OBJECTIVE NO. 2 JACKPILE-PAGUATE URANIUM MINE MEDIA OF CONCERN: SURFACE WATER/SEDIMENT

STEP 6. SPECIFY LIMITS ON DECISION ERRORS ((Continued)
ESTABLISH THE TRUE STATE OF NATURE FOR EACH DECISION RULE.	The true state of nature when the surface water/sediment is decided to be below three times the maximum background concentrations when in fact, it is not below three times the maximum background concentrations, is that the surface water/sediment does need remedial action. The true state of nature when the surface water/sediment is decided to be above three times the maximum background concentrations when in fact, it is not above three times the maximum background concentrations, is that the surface water/sediment does not need remedial action.
DEFINE THE TRUE STATE OF NATURE FOR THE MORE SEVERE DECISION ERROR AS THE BASELINE CONDITION OR THE NULL HYPOTHESIS (H ₀) AND DEFINE THE TRUE STATE FOR THE LESS SEVERE DECISION ERROR AS THE ALTERNATIVE HYPOTHESIS (H _a).	Ho: The surface water/sediment represented by the sample is above three times the maximum background concentrations. Ha: The surface water/sediment represented by the sample is below three times the maximum background concentrations.
ASSIGN THE TERMS "FALSE POSITIVE" AND "FALSE NEGATIVE" TO THE PROPER DECISION ERRORS.	 False Positive Error = Type I False Negative Error = Type II
ASSIGN PROBABILITY VALUES TO POINTS ABOVE AND BELOW THE ACTION LEVEL THAT REFLECT THE ACCEPTABLE PROBABILITY FOR THE OCCURRENCES OF DECISION ERRORS.	The assignment of probability values is not applicable to these DQOs because a nonprobablistic (judgment-based) process has been specified.
STEP 7. OPTIMIZE THE DESIGN	
REVIEW THE DQOs.	Surface water/sediment sample locations were selected based on the location of on-site waste source areas and proximity of the probable point of entry (PPEs) along the surface water pathway.

DEVELOP GENERAL SAMPLING AND ANALYSIS DESIGN.

Up to ten surface water and sediment samples will be collected from locations along the surface water pathway. The samples will be analyzed for CoCs to determine the presence of hazardous substances as defined by CERCLA, and to show potential attribution to the pathways of concern used for HRS evaluation. The samples will be analyzed for uranium isotopes, TAL metals, gross alpha, beta, and gamma, and radium. In addition, water samples will be analyzed for general cation/anion chemistry and total dissolved solids.

APPENDIX C

WESTON AND ERT STANDARD OPERATING PROCEDURES

SOP	1001.01					
GROUP	Sampling Procedur	Sampling Procedures				
SUB-GROUP	Soil Sampling Proc	Soil Sampling Procedures				
TITLE	Surface Soil Sampling					
DATE	11/19/2001	FILE	1001-01.DOC	PAGE	1 of 3	

INTRODUCTION

The following Standard Operating Procedure (SOP) is to describe the procedures for collecting representative soil samples. Analysis of soil samples may determine whether concentrations of specific soil pollutants exceed established action levels, or if the concentrations of soil pollutants present a risk to public health, welfare, or the environment. This SOP is similar to SOP Number 1001.03 for collecting near surface soil samples with a hand auger.

PROCEDURE

Surface soil samples may be collected using a variety of methods and equipment. The methods and equipment used are dependent on the depth of the desired sample, the type of sample required (disturbed versus undisturbed), and the type of soil. Near-surface soils may be easily sampled using a spade, trowel, or hand scoop.

Sample Preservation

Cooling to $4^{\circ}\text{C} \pm 2^{\circ}\text{C}$, supplemented by a minimal holding time, is suggested.

Interferences and Potential Problems

There are two primary interferences or potential problems associated with soil sampling: cross-contamination of samples and improper sample collection. Cross-contamination problems can be eliminated or minimized through the use of dedicated (disposable) sampling equipment. If this is not possible or practical, then decontamination of sampling equipment is necessary. Improper sample collection can involve using contaminated equipment, disturbance of the matrix resulting in compaction of the sample, or inadequate homogenization of the samples where required, resulting in variable, non-representative results. Homogenization may also affect sample representativeness where the analytical requirements include volatile organic compounds.

Equipment or Apparatus

The equipment used for sampling may be selected from the following list, as appropriate:

- Tape measure
- Survey stakes or flags
- Stainless steel, plastic, or other appropriate homogenization bucket or bowl
- Ziploc plastic bags
- Logbook
- Labels
- Chain-of-custody forms and seals
- Coolers
- Ice
- Decontamination supplies and equipment
- Canvas or plastic sheet
- Spatulas/spades/shovels
- Scoops

SOP	1001.01				
GROUP	Sampling Procedur	res			
SUB-GROUP	Soil Sampling Proc	edures			
TITLE	Surface Soil Sampl	ing			
DATE	11/19/2001	FILE	1001-01.DOC	PAGE	2 of 3

- Plastic or stainless steel spoons
- Trowel

Preparation

- 1. Determine the extent of the sampling effort, the sampling methods to be employed, and what equipment and supplies are required.
- 2. Obtain necessary sampling and monitoring equipment from the list above.
- 3. Prepare schedules, and coordinate with staff, client, and regulatory agencies, if appropriate.
- 4. Perform a general site survey prior to site entry in accordance with the site-specific health and safety plan.
- 5. Decontaminate or preclean equipment, and ensure that it is in working order.
- 6. Use stakes, buoys, or flagging to identify and mark all sampling locations. Consider specific site factors, including extent and nature of contaminant, when selecting sample locations. If required, the proposed locations may be adjusted based on site access, property boundaries, and surface obstructions. All staked locations will be utility-cleared by the property owner or other responsible party prior to soil sampling.
- 7. Evaluate safety concerns associated with sampling that may require use of personal protective equipment and/or air monitoring.

Surface Soil Sample Collection

Collect samples from the near-surface soil with tools such as spades, shovels, and scoops. Surface material can be removed to the required depth with this equipment, then a stainless steel or plastic scoop can be used to collect the sample. The use of a flat, pointed mason trowel to cut a block of the desired soil can be helpful when undisturbed profiles are required. A stainless steel scoop, lab spoon, or plastic spoon will suffice in most other applications. Avoid the use of devices plated with chrome or other target analyte materials.

The following procedures should be followed when collecting surface soil samples:

- 1. Carefully remove the top layer of soil or debris to the desired sample depth with a precleaned spade.
- 2. Using a pre-cleaned, stainless steel scoop, plastic spoon, or trowel, remove and discard a thin layer of soil from the area which came in contact with the spade.
- 3. If volatile organic analysis is to be performed, transfer a portion of the sample directly into an appropriate, labeled sample container(s) with a stainless steel lab spoon, plastic lab spoon, or equivalent and secure the cap(s) tightly. Place the remainder of the sample into a stainless steel, plastic, or other appropriate homogenization container, and mix thoroughly to obtain a homogenous sample representative of the entire sampling interval. Then, either place the sample into an appropriate, labeled container(s) and secure the cap(s) tightly; or if composite samples are to be collected, place a sample from another sampling interval into the

SOP	1001.01				
GROUP	Sampling Procedur	res			
SUB-GROUP	Soil Sampling Procedures				
TITLE	Surface Soil Samp	ing			
DATE	11/19/2001	FILE	1001-01.DOC	PAGE	3 of 3

homogenization container and mix thoroughly. When compositing is complete, place the sample into appropriate, labeled container(s) and secure the cap(s) tightly.

- 4. Fill hole created through sampling with unused material or other appropriate backfill material (sand).
- 5. Record applicable information into field log book or appropriate forms as documentation of sampling.

SOP	1001.04				
GROUP	Sampling Procedure	es			
SUB-GROUP	Soil Sampling Prod	ledures			
TITLE	Sampling of Stockp	oiled Soil			
DATE	11/19/2001	FILE	1001-04.DOC	PAGE	1 of 1

INTRODUCTION

The following Standard Operating Procedure (SOP) describes the procedure for collecting representative samples of stockpiled soil. Representative soil samples may be collected for analysis to determine whether concentrations of specific soil pollutants exceed established action levels, or if the concentrations of soil pollutants present a risk to public health and welfare, or the environment. Soil samples are also typically collected from stockpiles for classification prior to entry into a soil treatment process or offsite disposal, or after treatment to verify the effectiveness of the treatment system. This soil sampling procedure is closely related to SOP Nos. 1001.01, 1001.03, and 1001.10 regarding soil sampling procedures.

PROCEDURE

Stockpiles will be sampled as follows:

- Soil may be collected from the surface of a stockpile using the surface soil sampling procedure (SOP 1001.01) or from deeper within the stockpile according to the hand augering procedure (SOP 1001.03) as appropriate to obtain the required soil material. The procedure to be used to physically collect soil samples from stockpiles are described in SOP Nos. 1001.01, 1001.03, and 1001.10 (soil compositing). Reference should be made to these SOPs for specific sampling equipment, procedures, and other general guidelines. Equipment that may be used as part of the soil compositing procedure is identified under SOP Nos. 1001.01 and 1001.03 where general soil sampling methods are described.
- Each project may have different stockpile sampling objectives and requirements. Therefore, the sampling of stockpiles should be addressed in a site-specific Sampling and Analysis Plan and the soil sampling implemented in accordance with this plan.
- Samples may be collected from discrete locations in a pile and submitted for laboratory analysis, as described in SOP Nos. 1001.01 and 1001.03. More typical is that several samples from a single stockpile will be collected and composited to prepare a single sample for laboratory analysis. Collecting composite samples from a stockpile is recommended and will generally be performed to better characterize the soil in the pile. The number of samples to collected from a stockpile and composite will depend on the size of the stockpile and the particular requirements of the project. Typically compositing for characterization purposes is on the order of 1 composite soil sample for every 50 cubic yards. Compositing will be performed in accordance with SOP 1001.10.

REFERENCES

SOP No. 1001.01 - Standard Operating Procedure, Surface Soil Sampling

SOP No. 1001.03 - Standard Operating Procedure, Shallow Subsurface and Near Surface Soil Sampling

SOP No. 1001.10 - Standard Operating Procedure, Soil Compositing

SOP	1001.10				
GROUP	Soil Sampling Prod	cedures			
SUB-GROUP					
TITLE	Soil Compositing		_		
DATE	2/23/2010	FILE	1001-10.DOC	PAGE	1 of 2

INTRODUCTION

The following Standard Operating Procedure (SOP) describes the procedure for compositing soil samples. Soil samples are typically collected for laboratory analysis, and sometimes it is necessary to composite (mix together) samples from several locations for one combined analysis at the laboratory. This soil sampling procedure is closely related to SOP Nos. 1001.01, 1001.03, and 1001.10 regarding soil sampling procedures. This procedure serves as an alternative method of sample preparation prior to placing the samples in containers, as described in the other named SOPs.

PROCEDURE

Equipment

Equipment that may be used as part of the soil compositing procedure is identified under SOP Nos. 1001.01 and 1001.03 where soil sampling methods are described. Specific equipment typically used during the compositing process after discrete samples are collected includes:

- Mixing bowls or buckets
- Scoops, spatulas, and knives
- Sample containers
- Personal protection clothing
- Plastic Sheeting
- Decontamination equipment and supplies

Method

The procedure to be used to physically collect soil samples are described in SOP Nos. 1001.01 and 1001.03. Reference should be made to these SOPs for specific sampling equipment, procedures, and other general guidelines. As soil samples are collected, the site-specific Sampling and Analysis Plan may required compositing (mixing together) of two or more samples to create a single sample that will be sent to the laboratory for analysis. When this is the case, the following compositing procedure will generally be used:

- The soil will be collected in general accordance with SOP 1001.01 or 1001.03, with the exception that samples from discrete locations will generally not be immediately placed into sample containers and an additional preparation step (i.e., compositing) will be performed.
- As they are collected, soil samples selected for compositing will be staged in a clean mixing bowl or mixing bucket until each sample to be included in the composite sample is obtained. Depending on site requirements and analytical procedures to be requested, it may be necessary to temporarily stage individual discrete-location samples within clean sample jars, aluminum foil, or other appropriate materials for the project. The method for sample staging should be specified in the site-specific sampling and analysis plan.

SOP	1001.10				
GROUP	Soil Sampling Prod	cedures			
SUB-GROUP					
TITLE	Soil Compositing				
DATE	2/23/2010	FILE	1001-10.DOC	PAGE	2 of 2

- For composite samples that will be analyzed for volatile organic compounds, an equal portion of soil will be removed directly from each discrete-location sample and placed into a final sample jar without homogenizing the soil.
- For analyses other than volatile organics, equal portions of soil will be removed from each discrete-location sample and placed in a clean mixing bowl. The equal portions of the samples will then be broken up and homogenized together using a scoop or spatula. Homogenization will generally continue until the discrete samples being combined are reasonably indistinquishable as indivual individual samples in the soil mixture. However, it is recognized that homogenization can be difficult for highly plastic clays. In this case, equal amounts of the the soil core of each clay sample will be cut into small, roughly cubical pieces using a stainless steel knife, and an equal numbers of pieces of each discrete sample will be placed into the bowl and homogenized to extent practical.
- The composited soil sample will be collected from the mixing bowl containing the individual homogenized samples after homogenization is performed. The composited sample will be collected using a stainless steel or disposable plastic scoop or similar tool. The sample will be placed in a clean sample container and then handled in accordance with soil sampling SOPs 1001.01 and 1001.03.

Variations on this procedure are allowable to accommodate different soil conditions and any site requirements specifically identified in the site-specific Sampling and Analysis Plan.

The number of discrete samples that may be composited into a single sample typically ranges from two to six. The number of discrete samples that may be composited for the project in question will be specified in the site-specific Sampling and Analysis Plan.

REFERENCES

SOP No. 1001.01 - Standard Operating Procedure, Surface Soil Sampling

SOP No. 1001.03 - Standard Operating Procedure, Shallow Subsurface and Near Surface Soil Sampling

SOP	1002.01				
GROUP	Sampling Procedur	es			
SUB-GROUP	Surface Water				
TITLE	Surface Water Sam	pling			
DATE	11/19/2001	FILE	1002-01.DOC	PAGE	1 of 3

INTRODUCTION

The following Standard Operating Procedure (SOP) is to describe the procedures for collecting representative surface water samples. Analysis of surface samples may determine whether concentrations of specific soil pollutants exceed established action levels, or if the concentrations of pollutants present a risk to public health, welfare, or the environment.

PROCEDURE

Surface water samples may be collected using a variety of methods and equipment. The methods and equipment used are usually dependent on the location of the body of water being sampled. Sampling can be performed by merely submerging the sample container, a weighted-bottle sampler with stopper, a bailer, or by pump assisted methods. Several types of pumps can be used for sampling depending on the objectives of sampling and the site conditions.

Sample Preservation

Samples are to be preserved in conformance with the site-specific Quality Assurance Project Plan, Sampling and Analysis Plan or work plan. In general these requirements include refrigeration to 4°C, addition of appropriate additives (HCl, H₂SO₄, NaOH) to adjust and fix pH, and a defined maximum holding time. If a site-specific plan is not available, the analytical laboratory should be consulted for the appropriate preservation procedures.

Interferences and Potential Problems

There are two primary interferences or potential problems associated with surface water sampling: cross-contamination of samples and improper sample collection. Cross-contamination problems can be eliminated or minimized through the use of dedicated sampling equipment. If this is not possible or practical, then decontamination of sampling equipment is necessary. Improper sample collection can involve using contaminated equipment, undue disturbance of the sample matrix, or improper sample location.

Equipment or Apparatus

- Ziploc plastic bags
- Logbook
- Labels
- Chain-of-custody forms and seals
- Coolers

- Ice
- Decontamination supplies and equipment
- Discharge tubing
- Sample containers
- Sampling devices

SOP	1002.01				
GROUP	Sampling Procedur	es			
SUB-GROUP	Surface Water	Surface Water			
TITLE	Surface Water Sam	pling			
DATE	11/19/2001	FILE	1002-01.DOC	PAGE	2 of 3

Preparation

- 1. Determine the extent of the sampling effort, the sampling methods to be employed, and which equipment and supplies are required.
- 2. Obtain necessary sampling and monitoring equipment.
- 3. Decontaminate or preclean equipment, and ensure that it is in working order.
- 4. Prepare schedules, and coordinate with staff, client, and regulatory agencies, if appropriate.
- 5. Perform a general site survey prior to site entry in accordance with the site-specific health and safety plan.

Surface Water Sampling

Samples from shallow depths can be readily collected by merely submerging the sample container. In flowing surface water bodies, the container's mouth should be positioned so that it faces upstream, while the sampling personnel stand downstream so as not to stir up sediment that could potentially contaminate the sample.

Collecting a representative sample from a larger body of surface water requires that samples be collected near the shore unless boats are feasible and permitted. If boats are used, the body of water should be cross sectioned, and samples should be collected at various depths across the body of water in accordance with the specified sampling plan. For this type of sampling, a weighted-bottle sampler is used to collect samples at a predetermined depth. The sampler consists of a glass bottle, a weighted sinker, a bottle stopper, and a line that is used to open the bottle and to lower and raise the sampler during sampling. The procedure for use is as follows:

- Assemble the weighted bottle sampler.
- Gently lower the sampler to the desired depth so as not to remove the stopper prematurely.
- Pull out the stopper with a sharp jerk of the sampler line.
- Allow the bottle to fill completely, as evidenced by the cessation of air bubbles.
- Raise the sampler and cap the bottle.
- Wipe the bottle clean. The sampling bottle can be also be used as the sample container for shipping.

Teflon bailers have also been used where feasible for collecting samples in deep bodies of water.

SOP	1002.01				
GROUP	Sampling Procedur	res			
SUB-GROUP	Surface Water	Surface Water			
TITLE	Surface Water Sam	Surface Water Sampling			
DATE	11/19/2001	FILE	1002-01.DOC	PAGE	3 of 3

Another method of extending the reach of sampling efforts is the use of a small peristaltic pump. In this method the sample is drawn through heavy-wall Teflon tubing and pumped directly into the sample container. This system allows the operator to reach into the liquid body, sample from depth, or sweep the width of narrow streams.

The general sampling procedures are listed below:

- 1. Collect the sample using whichever technique, submerged bottle, bottle sampler with stopper, pump & tubing, or bailer.
- 2. The collected sample may be collected in the sample containers or may be transferred to the appropriate sample containers in order of the volatile organics first and inorganics last.
- 3. Label sample containers, place on ice in a cooler, remove, and decontaminate equipment as necessary.

REFERENCES

SOP 0110.01	Sample Nomenclature
SOP 1005.01	Field Duplicate Collection
SOP 1005.02	Rinse Blank Preparation
SOP 1005.03	Field Blank Preparation
SOP 1101.01	Sample Custody - Field
SOP 1102.01	Sample Shipping
SOP 1201.01	Sampling Equipment Decontamination
SOP 1501.01	Field Logbook

SOP	1002.02				
GROUP	Sampling Procedur	es			
SUB-GROUP	Water Sampling Pr	Water Sampling Procedures			
TITLE	Groundwater Sampling				
DATE	11/19/2001	FILE	1002-02.DOC	PAGE	1 of 3

The following Standard Operating Procedure (SOP) is to describe the procedures for collecting representative groundwater samples. Analysis of groundwater samples may determine whether concentrations of specific pollutants exceed established action levels, or if the concentrations of pollutants present a risk to public health, welfare, or the environment. This SOP applies to collection of groundwater samples from monitoring wells and/or piezometers.

PROCEDURE

Groundwater samples may be collected using a variety of methods and equipment. The methods and equipment used are dependent on the depth of the piezometric level from the ground surface and the particular analytes of concern. For example, deep monitoring wells are typically purged and sampled using submersible pumps and shallow monitoring wells are typically purged and sampled using bailing techniques.

Sample Preservation

Preservation depends on the particular analyte or analyte group as specified in EPA approved analytical methods. In almost all cases, cooling to $4^{\circ}\text{C} \pm 2^{\circ}\text{C}$, supplemented by a minimal holding time is required.

Interferences and Potential Problems

There are two primary interferences or potential problems associated with groundwater sampling: cross-contamination of samples and improper sample collection. Cross-contamination problems can be eliminated or minimized through the use of dedicated (disposable) sampling equipment. If this is not possible or practical, then decontamination of sampling equipment is necessary. Improper sample collection can involve using contaminated equipment, inadequate purging of static and potentially unrepresentative water from the well, improper purging and sampling methods (creating turbidity problems), and improper handling of samples allowing atmospheric alteration and non-representative results (i.e. volatilization).

Equipment or Apparatus

The equipment used for sampling may be selected from the following list, as appropriate:

- Water level indicator
- Drums or buckets as appropriate to contain purge water
- Ziploc plastic bags
- Logbook
- Labels
- Chain-of-custody forms and seals
- Coolers
- Ice
- Decontamination supplies and equipment
- Canvas or plastic sheet

SOP	1002.02				
GROUP	Sampling Procedur	es			
SUB-GROUP	Water Sampling Pr	ocedures			
TITLE	Groundwater Sampling				
DATE	11/19/2001	FILE	1002-02.DOC	PAGE	2 of 3

- Physical monitoring equipment (pH, specific conductivity, temperature, redox potential, and dissolved oxygen measuring devices)
- Bailers (disposable and non-disposable) and bailer rope
- Submersible or peristaltic pumps and associated tubing
- Power supply as appropriate for the selected pump
- Sample containers

Preparation

- 1. Determine the purpose of the sampling effort, the sampling methods to be employed, and what equipment and supplies are required.
- 2. Obtain necessary sampling and monitoring equipment from the list above.
- 3. Prepare schedules, and coordinate with staff, client, and regulatory agencies, if appropriate.
- 4. Perform a general site survey prior to site entry in accordance with the site-specific health and safety plan.
- 5. Decontaminate or preclean equipment, and ensure that it is in working order.
- 6. Identify the monitoring wells to be sampled. Consider specific site factors, including extent and nature of contaminant, when selecting sample locations.
- 7. Evaluate safety concerns associated with sampling that may require use of personal protective equipment and/or air monitoring.

Groundwater Sample Collection

For newly installed monitoring wells and piezometers, sampling should not be performed sooner than 24 hours after development. Prior to sampling, each well/piezometer will be purged to remove static groundwater and allow collection of representative samples. Purging can be performed by either pumping with a submersible pump or peristaltic pump or by bailing with a bailer of appropriate construction. Sampling can be performed directly by whichever method is used. If a submersible pump is used as the sampling device, it will be a variable discharge type, and sampling will be performed at low flow to avoid induced volatilization of constituents (if present) from the impellers. The following procedure will be used for monitoring well/piezometer sampling.

- 1. Determine the construction details (screen interval, size of borehole, length of filter pack, etc.) of each monitoring well to be sampled. This information can be obtained from well construction diagrams and/or boring logs.
- 2. Measure and record the static water level and total depth of the well from the top of the well casing (or identified reference point). An oil/water interphase probe will be used to determine the water level and presence of phase-separated product, if present. If phase separated product is present in the well (either NAPL or DNAPL) an evaluation will be performed as to the necessity of sampling either the well or product will be performed. In the event that accumulated sediment is identified in the wells, redevelopment may be necessary prior to sampling.
- 3. Determine the volume of water to be purged from the well/piezometer.

SOP	1002.02				
GROUP	Sampling Procedure	es			
SUB-GROUP	Water Sampling Pro	Water Sampling Procedures			
TITLE	Groundwater Samp	ling			
DATE	11/19/2001	FILE	1002-02.DOC	PAGE	3 of 3

- 4. Lay plastic sheeting or canvas around the well to prevent contact of the bailer line or pump cable and tubing with the ground.
- 5. Evacuate a minimum of three well volumes of water from the well/piezometer. Low flow purging is recommended to reduce agitation and turbidity. In general, three wetted casing plus annular volumes of water are required for successful purging. For instances of low yield, purging to dryness twice is acceptable. Measure and record pH, specific conductivity, temperature and any other indicators of interest throughout purging activities. Purging is considered complete when the volume requirements are satisfied, the physical measurements stabilize (typically 0.5 pH units, 10% specific conductivity, and 1° C temperature), and when the evacuated water is reasonably free of sediment.
- 6. Allow the well to recover (ideally 75% of the static level), and collect the appropriate groundwater samples. A variable discharge submersible pump, peristaltic pump, or disposable bailer can be used to collect the groundwater samples. All groundwater samples will be collected and transferred to the appropriate labeled sample containers, preserved as necessary, and placed in a cooler with ice.
- 7. Record applicable information into field log book or appropriate forms as documentation of sampling.
- 8. Properly package and ship samples to the designated laboratory as soon as possible (typically the same day as collected) following proper chain-of-custody and shipping procedures.

SOP	1002.04				
GROUP	Sampling Procedur	es			
SUB-GROUP	Soil Sampling Proc	Soil Sampling Procedures			
TITLE	Sediment Sampling				
DATE	2/23/2010	FILE	1002-04.DOC	PAGE	1 of 3

The following Standard Operating Procedure (SOP) is to describe the procedures for collecting representative sediment samples using a trowel, piston corer, WILDCO KB Core Sampler, a Ponar Grab Sampler, or other similar equipment. Analysis of sediment samples may be performed to determine whether concentrations of specific sediment pollutants exceed established action levels, or if the concentrations of sediment pollutants present a risk to public health, welfare, or the environment.

PROCEDURE

Overview

Sediment samples may be collected using trowels, core and Ponar sampler, or a variety of similar methods and equipment. The methods and equipment used are dependent on the depth of the desired sample, the type of sample required (disturbed versus undisturbed), and the type of sediment (fines versus coarse). Sampling in shallow areas or streams near the surface may only require a hand trowel, while sampling at depth may be performed using a core or Ponar sampler.

Sample Preservation

Refrigeration to 4° C \pm 2° C, supplemented by a minimal holding time, is suggested.

Interferences and Potential Problems

There are two primary interferences or potential problems associated with sediment sampling: cross-contamination of samples and improper sample collection. Cross-contamination problems can be eliminated or minimized through the use of dedicated (disposable) sampling equipment. If this is not possible or practical, then decontamination of sampling equipment is necessary. Improper sample collection can involve using contaminated equipment, disturbance of the matrix resulting in mixing of the sample, or inadequate homogenization of the samples where required, resulting in variable, non-representative results. Homogenization may also affect sample representativeness when the analytical requirements include volatile organic compounds.

Equipment or Apparatus

The equipment selected for the sampling effort may include the following as appropriate:

- Tape measure
- Survey stakes or flags
- Stainless steel, plastic, or other appropriate homogenization bucket or bowl
- Ziploc plastic bags
- Logbook
- Labels
- Chain-of-custody forms and seals
- Coolers
- Ice

SOP	1002.04				
GROUP	Sampling Procedur	es			
SUB-GROUP	Soil Sampling Proc	Soil Sampling Procedures			
TITLE	Sediment Sampling				
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- Decontamination supplies and equipment (i.e. brushes and buckets)
- Canvas or plastic sheeting
- Spatulas
- Scoops
- Plastic or stainless steel spoons
- Trowel
- Auger bucket
- Extension rods
- T-handle
- KB Core Sampler
- Ponar Grab Sampler
- Air monitor

Preparation

- 1. Determine the extent of the sampling effort, the sampling methods to be employed, and which equipment and supplies are required.
- 2. Obtain necessary sampling and monitoring equipment from the list above. Additional equipment may be added to this list as appropriate to perform other sampling.
- 3. Decontaminate or preclean equipment, and ensure that it is in working order.
- 4. Perform a general site survey prior to site entry in accordance with the site-specific health and safety plan.
- 5. Use stakes, buoys, or flagging to identify and mark all sampling locations. Consider specific site factors, including extent and nature of contaminant, when selecting sample locations. If required, the proposed locations may be adjusted based on site access, property boundaries, and obstructions.

Sediment Sampling in Shallow Waters

The following procedures should be used when collecting sediment samples in shallow waters:

- 1. Collect sediments as specified in the work plan or as determined during office preparation activities, using a stainless steel trowel, piston corer or similar device and a stainless steel, tempered glass or aluminum container.
- 2. Standing downstream of the sample stations, collect discrete sediment samples from each station and, if required in the work plan, composite in stainless steel, tempered glass or aluminum container.
- 3. Collect sediment samples of deposited material from the depth specified in the work plan or as determined during the office preparation activities. Record the depth in the logbook. Selective removal of the top sediment layers may be required and should be accomplished by carefully removing the sediments with a stainless steel trowel or scoop. In streams where water velocity is insufficient to disturb sediment fines during sediment sampling, a stainless

SOP	1002.04					
GROUP	Sampling Procedur	es				
SUB-GROUP	Soil Sampling Proc	Soil Sampling Procedures				
TITLE	Sediment Sampling	7				
DATE	2/23/2010	FILE	1002-04.DOC	PAGE	3 of 3	

steel trowel or scoop may be used for sampling. Where water velocities are high, a stainless steel corer will be utilized.

- 4. When applicable, composite discrete sediment samples by placing equal volumes of sediment material collected from the sample points into the container and mixing thoroughly to obtain a homogeneous mixture. Samples may be sieved or hand picked, if necessary, to remove larger materials, such as leaves, sticks, gravel, or rocks. Record in the logbook the nature of any materials removed from the sediment samples.
- 5. Place each sediment sample into the proper clean, unused sample container, as required by the work plan or laboratory. Sampling personnel must avoid placing sediment into the sample container and decanting off the excess liquid in analyzing for volatile organics and water soluble compounds in the sediment and reduces accurate representation of sediment analysis.
- 6. Fill out labels with waterproof ink and attach to the sample container.
- 7. Decontaminate sampling equipment between samples.

Sediment Sampling in Deep Waters

Procedures for sampling in deep waters are the same as for shallow waters except the sampling equipment is different. Soft, fine-grained sediments collected in deep waters will be sampled with a WILDCO KB Core Sampler or similar equipment . Coarse-grained sediments will be collected utilizing a Ponar Grab Sampler or similar equipment. Both samplers will be operated from a boat following appropriate safety procedures. Documentation, containerization, labeling and decontamination procedures are the same as for sediment samples collected in shallow waters.

Sediment Sampling in Drainage Ditches and Intermittent Streams

Procedures for sediment sampling in drainage ditches and the dry portions of intermittent streams are as specified for shallow water sediments.

SOP	1005.01				
GROUP	Sampling Procedur	es			
SUB-GROUP	Field QA/QC Sam	Field QA/QC Sampling			
TITLE	Field Duplicate Collection				
DATE	4/27/2005	FILE	1005-01.DOC	PAGE	1 of 2

The following Standard Operating Procedure (SOP) describes the procedure for collecting field duplicate soil and water samples. When samples are collected for analysis, it is typically desired that independent data allowing evaluation of laboratory precision (i.e., the degree to which a laboratory result can be repeated) on site-specific samples be collected.

A field duplicate sample is a second sample collected at the same location as the original sample. Duplicate samples are collected simultaneously or in immediate succession, using identical recovery techniques, and treated in an identical manner during storage, transportation, and analysis. The sample containers are assigned an identification number in the field such that they cannot be identified (blind duplicate) as duplicated samples by laboratory personnel performing the analysis. Specific locations are designated for collection of field duplicate samples prior to the beginning of sample collection.

The duplicate soil sampling procedure is closely related to SOP Nos. 1001.01, 1001.03, and 1001.10 regarding soil sampling procedures. This procedure serves as an alternative method or extension of sample preparation prior to placing the samples in containers, as described in the 1001 series of the SOPs (e.g. 1001.01 and 1001.03).

DUPLICATE SOIL SAMPLING PROCEDURE

The procedure to be used to physically collect soil samples are described in SOP Nos. 1001.01 and 1001.03. Reference should be made to these SOPs for specific sampling equipment, procedures, and other general guidelines. As soil is collected, the following procedure will be used to prepare a field duplicate sample:

- The soil will be collected in general accordance with SOP 1001.01 or 1001.03, with the exception that samples will generally not be immediately placed into sample containers and an additional preparation step (i.e., sample splitting) will be performed.
- As they are collected, soil samples to be submitted as field duplicates will be staged in a clean mixing bowl or mixing bucket.
- For samples that will be analyzed for volatile organic compounds, the soil sample will be split in half and an equal portion of soil will be placed directly into two or more different sample containers, each container representing a different sample for laboratory analysis. The soil will not be homogenized to minimize the potential for volatilization of the organic compounds potentially in the sample.
- For analyses of chemicals other than volatile organic compounds, the soil removed from the discrete sample location will be homogenized in a clean mixing bowl using a clean scoop or spatula (as described in SOPs 1001.01 and 1001.03). Homogenization will generally continue until the discrete samples being combined are reasonably indistinguishable as individual samples in the soil mixture. However, it is recognized that homogenization can be difficult for highly plastic clays. In this case, equal amounts of the soil core of each clay sample will be cut into small, roughly cubical pieces using a stainless steel knife and placed into a bowl and homogenized to extent practical.

SOP	1005.01				
GROUP	Sampling Procedur	es			
SUB-GROUP	Field QA/QC Sam	Field QA/QC Sampling			
TITLE	Field Duplicate Co	Field Duplicate Collection			
DATE	4/27/2005	FILE	1005-01.DOC	PAGE	2 of 2

• The field duplicate sample (except for volatiles as note above) will be collected from the mixing bowl containing the homogenized samples after homogenization is performed. The composited sample will be collected using a stainless steel or disposable plastic scoop or similar tool. The sample will be placed in a clean sample container and then handled in accordance with soil sampling SOPs 1001.01 and 1001.03.

Another difference from the referenced SOPs is that additional soil volume may need to be collected from a discrete sample location during the sampling process to provide sufficient sample volume for two or more sets of laboratory analyses. If the collection of additional sample volume will result in the sample interval expanding to greater depths or laterally outward, the sampling tools identified in 1001 series of the SOPs can be used at two immediately vertically or laterally adjacent locations, as appropriate. If sampling from two adjacent but distinct locations is necessary to obtain adequate sample volume, the soil from the two locations should be composited in accordance with SOP 1001.10. Field duplicates of composited samples may also be performed using this SOP for field duplicate samples.

Variations on this procedure are allowable to accommodate different soil conditions and any site requirements specifically identified in the site-specific Sampling and Analysis Plan. Equipment that may be used as part of the soil compositing procedure is identified under SOP Nos. 1001.01 and 1001.03 where soil sampling methods are described.

DUPLICATE WATER SAMPLING PROCEDURES

The procedure to be used to physically collect water samples are described in 1002 series of the SOPs (e.g. 1002.01 and 1002.02). Reference should be made to these SOPs for specific sampling equipment, procedures, and other general guidelines. A duplicate water sample will be collected from the same location as the parent sample and within 15 minutes of the collection of the parent sample.

The number of samples that may be submitted as blind field duplicates for the project in question will be specified in the site-specific sampling plan. Blind field duplicates are typically collected at a frequency of 1 per 10 samples of a given environmental media at sites, especially where laboratory analytical data will be used for evaluating regulatory compliance and other engineering judgments. Sampling in support of a routine monitoring program may not require field duplicates. Reference should be made to the site-specific contract and work plans.

REFERENCES

SOP No. 1001.01 - Standard Operating Procedure, Surface Soil Sampling

SOP No. 1001.03 - Standard Operating Procedure, Soil Sampling - Hand Auger Method

SOP No. 1001.10 - Standard Operating Procedure, Soil Compositing

SOP	1005.02				
GROUP	Sampling Procedur	es			
SUB-GROUP	Field QA/QC Sam	Field QA/QC Sampling			
TITLE	Rinse Blank Preparation				
DATE	2/6/2009	FILE	1005-02.DOC	PAGE	1 of 1

The following Standard Operating Procedure (SOP) presents a method to prepare a type of quality control sample specific to the field decontamination process, the equipment rinse blank. The rinse blank provides information on the effectiveness of the decontamination process employed in the field. When used in conjunction with field blanks and trip blanks, the rinse blank can be used to assist in evaluating possible compromise of samples from field related activities.

PROCEDURE

The equipment rinse blank is prepared by passing target analyte-free (i.e., deionized) water over and through a field decontaminated sampling device, then collecting the rinse water in appropriate clean sample containers. Rinse blanks will typically be collected from equipment that comes in contact with samples, such as auger buckets, split spoons, bailers, shelby tubes, and stainless steel spoons/trowels. The collected sample will be coded appropriately prior to logging and shipping. Equipment blanks are not required if dedicated sampling equipment is used. Equipment blanks will be collected periodically during the day immediately after decontamination of the sampling equipment being used.

The frequency for collecting equipment blanks will be determined prior to engaging in field activities, and communicated in site-specific quality assurance project plans, sampling and analyses plans, or a type of work plan. Equipment blanks will be collected at a rate relative to each type of sample collection procedure (i.e., surface sample, sample at depth using a hand auger). Equipment blanks will generally be collected at a frequency of 1 per 20 (normal) samples of a given matrix.

SOP	1101.01				
GROUP	Sampling Handling	7			
SUB-GROUP	Sample Custody				
TITLE	Sample Custody in	Sample Custody in the Field			
DATE	11/19/2001	FILE	1101-01.DOC	PAGE	1 of 4

The following Standard Operating Procedure (SOP) presents procedures for maintaining sample chain of custody (COC) during activities where samples are collected.

PROCEDURE

Sample custody is defined as being under a person's custody if any of the following conditions exist:

- it is in their possession,
- it is in their view, after being in their possession,
- it was in their possession and they locked it up, or
- it is in a designated secure area.

A designated field sampler will be personally responsible for the care and custody of collected samples until they are transferred to another person or properly dispatched to the laboratory. To the extent practicable, as few people as possible will handle the samples.

Sample tags or labels will be completed and applied to the container of each sample. When the tags or labels are being completed, waterproof ink will be used. If waterproof ink is not used, the tags or labels will be covered by transparent waterproof tape. Sample containers may also be placed in Ziploc-type storage bags to help keep them clean in the cooler. Information typically included on the sample tags or labels will include the following:

- Project Code
- Station Number and Location
- Sample Identification Number
- Date and Time of Sample Collection
- Type of Laboratory Analysis Required
- Preservation Required, if applicable
- Collector's Signature
- Priority (optional)
- Other Remarks

Additional information may include:

- Anticipated Range of Results (Low, Medium, or High)
- Sample Analysis Priority

SOP	1101.01				
GROUP	Sampling Handling	5			
SUB-GROUP	Sample Custody				
TITLE	Sample Custody in	Sample Custody in the Field			
DATE	11/19/2001	FILE	1101-01.DOC	PAGE	2 of 4

A COC form will be completed each time a sample or group of samples is prepared for transfer to the laboratory. The form will repeat the information on each of the sample labels and will serve as documentation of handling during shipment. The minimum information requirements of the COC form are listed in Table 1101.01-A. An example COC form is shown in Figure 1101.01-A. The completed COC must be reviewed by the Field Team Leader or Site Manager prior to sample shipment. The COC form will remain each sample shipping container at all times, and another copy will be retained by the member of the sampling team who originally relinquished the samples or in a project file.

SOP	1101.01				
GROUP	Sampling Handling	7			
SUB-GROUP	Sample Custody				
TITLE	Sample Custody in	the Field			
DATE	11/19/2001	FILE	1101-01.DOC	PAGE	3 of 4

TABLE 1101.01-A CHAIN OF CUSTODY FORM

INFORMATION	COMPLETED BY	DESCRIPTION
COC	Laboratory	enter a unique number for each chain of custody form
SHIP TO	Field Team	enter the laboratory name and address
CARRIER	Field Team	enter the name of the transporter (e.g., FedEx) or handcarried
Airbill	Field Team	enter the airbill number or transporter tracking number (if applicable)
PROJECT NAME	Field Team	enter the project name
SAMPLER NAME	Field Team	enter the name of the person collecting the samples
SAMPLER SIGNATURE	Field Team	signature of the person collecting the samples
SEND RESULTS TO	Field Team	enter the name and address of the prime contractor
FIELD SAMPLE ID	Field Team	enter the unique identifying number given to the field sample (includes MS, MSD, field duplicate and field blanks)
DATE	Field Team	enter the year and date the sample was collected in the format M/D (e.g., 6/3)
Тіме	Field Team	enter the time the sample was collected in 24 hour format (e.g., 0900)
MATRIX	Field Team	enter the sample matrix (e.g., water, soil)
Preservative	Field Team	enter the preservative used (e.g., HNO3) or "none"
Filtered/ Unfiltered	Field Team	enter "F" if the sample was filtered or "U" if the sample was not filtered
Containers	Field Team	enter the number of containers associated with the sample
MS/MSD	Field Team or Laboratory	enter "X" if the sample is designated for the MS/MSD
ANALYSES REQUESTED	Field Team	enter the method name of the analysis requested (e.g., SW6010A)
COMMENTS	Field Team	enter comments
SAMPLE CONDITION UPON RECEIPT AT LABORATORY	Laboratory	enter any problems with the condition of any sample(s)
Cooler Temperature	Laboratory	enter the internal temperature of the cooler, in degrees C, upon opening
SPECIAL INSTRUCTIONS/COMME NTS	Laboratory	enter any special instructions or comments
RELEASED BY (SIG)	Field Team and Laboratory	enter the signature of the person releasing custody of the samples
COMPANY NAME	Field Team and Laboratory	enter the company name employing the person releasing/receiving custody
RECEIVED BY (SIG)	Field Team and Laboratory	enter the signature of the person receiving custody of the samples
DATE	Field Team and Laboratory	enter the date in the format M/D/YY (e.g., 6/3/96) when the samples were released/received
Тіме	Field Team and Laboratory	enter the date in 24 hour format (e.g., 0900) when the samples were released/received

SOP	1101.01				
GROUP	Sampling Handling	7			
SUB-GROUP	Sample Custody				
TITLE	Sample Custody in	Sample Custody in the Field			
DATE	11/19/2001	FILE	1101-01.DOC	PAGE	4 of 4

FIGURE 1101.01-A CHAIN OF CUSTODY FORM

SOP	1102.01				
GROUP	Sample Handling				
SUB-GROUP	Sample Shipping				
TITLE	Sample Shipping				
DATE	11/19/2001	FILE	1102-01.DOC	PAGE	1 of 1

The following Standard Operating Procedure (SOP) presents the procedures for sample shipping that will be implemented during field work involving sampling activities.

TERMS

COC - Chain-of-Custody

PROCEDURE

Prior to shipping or transferring custody of samples, they will be packed according to D.O.T. requirements with sufficient ice to maintain an internal temperature of $4^{\circ}C \pm 2^{\circ}C$ during transport to the laboratory. Samples relinquished to the participating laboratories will be subject to the following procedures for transfer of custody and shipment:

- 1. Samples will be accompanied by a COC record. When transferring possession of samples, the individuals relinquishing and receiving the samples will sign, date, and note the time of the sample transfer on the record. If sent by common carrier, a bill of lading or airbill should be used. Bill of lading and airbill receipts will be retained in the project file as part of the permanent documentation of sample shipping and transfer. This custody record documents transfer of sample custody from the sampler to another person or to the laboratory. The designated laboratory will accept custody in the field upon sample pick-up or at the laboratory if the samples are delivered via field personnel or a courier service.
- 2. Samples will be properly packed in approved shipping containers for laboratory pick-up by the appropriate laboratory for analysis, with separate, signed custody records enclosed in each sample box or cooler. Sample shipping containers will be padlocked or custody-sealed for transfer to the laboratory. The preferred procedure includes use of a custody seal wrapped across filament tape that is wrapped around the package at least twice. The custody seal will then be folded over and stuck to itself so that the only access to the package is by cutting the filament tape or breaking the seal to unwrap the tape. The seal will then be signed. The designated laboratory will accept custody of the samples upon receipt.
- 3. Whenever samples are split with state representatives or other parties, the COC record will be marked to indicate with whom the samples were split.
- 4. The field sampler will call the designated laboratory to inform them of sample shipment and verify sample receipt as necessary.

SOP	1201.01					
GROUP	Decontamination					
SUB-GROUP	Sampling Equipme	nt Decontamination				
TITLE	Sampling Equipme	Sampling Equipment Decontamination				
DATE	11/19/2001	FILE	1201-01.DOC	PAGE	1 of 3	

The following Standard Operating Procedure (SOP) presents the methods used for minimizing the potential for cross-contamination, and provides general guidelines for sampling equipment decontamination procedures.

PROCEDURE

As part of the Health and Safety Plan (HASP), develop and set up a decontamination plan before any personnel or equipment enter the areas of potential exposure. The decontamination plan should include the following:

- The number, location, and layout of decontamination stations
- Which decontamination apparatus is needed
- The appropriate decontamination methods
- Methods for disposal of contaminated clothing, apparatus, and solutions

Decontamination Methods

Personnel, samples, and equipment leaving the contaminated area of a site will be decontaminated. Various decontamination methods will be used to either physically remove contaminants, inactivate contaminants by disinfection or sterilization, or both. The physical decontamination techniques appropriate for equipment decontamination can be grouped into two categories: abrasive methods and non-abrasive methods.

Abrasive Cleaning Methods

Abrasive cleaning methods work by rubbing/scrubbing the surface containing the contaminant. This method includes mechanical and wet blasting methods.

Mechanical cleaning methods use brushes of metal or nylon. The amount and type of contaminants removed will vary with the hardness of bristles, length of brushing time, and degree of brush contact.

Cleaning can also be accomplished by water blasting which is also referred to as steam cleaning and pressure washing. Pressure washing utilizes high-pressure that is sprayed from a nozzle onto sampling equipment to physically remove soil or (potentially) contaminated material. Steam cleaning is a modification of pressure washing where the water is heated to temperatures approaching 100 °C to assist in removing organic constituents from equipment.

SOP	1201.01				
GROUP	Decontamination				
SUB-GROUP	Sampling Equipme	Sampling Equipment Decontamination			
TITLE	Sampling Equipme	Sampling Equipment Decontamination			
DATE	11/19/2001	FILE	1201-01.DOC	PAGE	2 of 3

Disinfection/Rinse Methods

Disinfectants are a practical means of inactivating chemicals or contaminants of concern. Standard sterilization methods involve heating the equipment which is impractical for large equipment. Rinsing removes contaminants through dilution, physical attraction, and solubilization.

The use of distilled/deionized water commonly available from commercial vendors may be acceptable for decontamination of sampling equipment provided that it has been verified by laboratory analysis to be target analyte free. Tap water may be used from any municipal water treatment system for mixing of decontamination solutions. An untreated potable water supply is not an acceptable substitute for tap water. Acids and solvents are occasionally utilized in decontamination of equipment to remove metals and organics, respectively, from sampling equipment. Other than ethanol, these are avoided when possible due to the safety, disposal, and transportation concerns associated with them.

Equipment or apparatuses that may be selected for use include the following:

- Personal protective clothing
- Non-phosphate detergent
- Selected solvents for removal of polar and nonpolar organics (ethanol, methanol, hexane)
- Acid washes for removal of metals (nitric acid)
- Long-handled brushes
- Drop cloths or plastic sheeting
- Paper towels
- Galvanized tubs or buckets
- Distilled, deionized, or tap water (as required by the project)
- Storage containers for spent wash solutions
- Sprayers (pressurized and non-pressurized)
- Trash bags
- Safety glasses or splash shield

Field Sampling Equipment Cleaning Procedures

The following procedures should be followed:

- 1. Where applicable, follow physical removal procedures previously described (pressure wash, scrub wash)
- 2. Wash equipment with a non-phosphate detergent solution
- 3. Rinse with tap water
- 4. Rinse with distilled or deionized water
- 5. Rinse with 10% nitric acid if the sample will be analyzed for metals/organics
- 6. Rinse with distilled or deionized water
- 7. Use a solvent rinse (pesticide grade) if the sample will be analyzed for organics
- 8. Air dry the equipment completely
- 9. Rinse again with distilled or deionized water

SOP	1201.01					
GROUP	Decontamination					
SUB-GROUP	Sampling Equipme	nt Decontamination				
TITLE	Sampling Equipme	Sampling Equipment Decontamination				
DATE	11/19/2001	FILE	1201-01.DOC	PAGE	3 of 3	

10. Place in clean bag or container for storage/transport to subsequent sampling locations.

Selection of the solvent for use in the decontamination process is based on the contaminants present at the site. Solvent rinses are not necessarily required when organics are not a contaminant of concern and may be eliminated from the sequence specified below. Similarly, an acid rinse is not required if the analyses do not include inorganics. Use of a solvent is required when organic contamination is present on-site. Typical solvents used for removal of organic contaminants include acetone, ethanol, hexane, methanol, or water. An acid rinse step is required if metals are present on-site. If a particular contaminant fraction is not present at the site, the tenstep decontamination procedure listed above may be modified for site specificity.

Sampling equipment that requires the use of plastic tubing should be disassembled and the tubing replaced with clean tubing before commencement of sampling and between sampling locations. Plastic tubing should not be reused.

SOP	1501.01				
GROUP	Field Documentation	Field Documentation			
SUB-GROUP					
TITLE	Field Logbook				
DATE	11/19/2001	FILE	1501-01.DOC	PAGE	1 of 3

The following Standard Operating Procedure (SOP) presents the procedures for documenting activities observed or completed in the field in a field logbook. The documentation should represent all activities of WESTON personnel and entities under WESTON's supervision.

TERMS

FSP - Field Sampling Plan

SAP - Sampling and Analysis Plan

QAPP - Quality Assurance Project Plan

HASP - Health and Safety Plan

PROCEDURE

Field logbooks will be used and maintained during field activities to document pertinent information observed or completed by WESTON personnel or entities that WESTON is responsible for providing oversight. Field logbooks are legal documents that form the basis for later written reports and may serve as evidence in legal proceedings. The Site Manager or Field Team Leader will review field log entries daily and initial each page of entries. Field logbooks will be maintained by the Site Manager or Field Team Leader during field activities and transferred to the project files for a record of activities at the conclusion of the project. General logbook entry procedures are listed below.

- Logbooks must be permanently bound with all pages numbered to the end of the book. Entries should begin on page 1.
- Only use blue or black ink (waterproof) for logbook entries.
- Sign entries at the end of the day, or before someone else writes in the logbook.
- If a complete page is not used, draw a line diagonally across the blank portion of the page and initial and date the bottom line.
- If a line on the page is not completely filled, draw a horizontal line through the blank portion.
- Ensure that the logbook clearly shows the sequence of the day's events.
- Do not write in the margins or between written lines, and do not leave blank pages to fill in later.
- If an error is made, make corrections by drawing a single line through the error and initialing it.
- Maintain control of the logbook and keep in a secure location.

SOP	1501.01				
GROUP	Field Documentation	Field Documentation			
SUB-GROUP					
TITLE	Field Logbook				
DATE	11/19/2001	FILE	1501-01.DOC	PAGE	2 of 3

Field logbooks will contain, at a minimum, the following information, if applicable:

General Information

- Name, location of site, and work order number
- Name of the Site Manager or Field Team Leader
- Names and responsibilities of all field team members using the logbook (or involved with activities for which entries are being made)
- Weather conditions
- Field observations
- Names of any site visitors including entities that they represent

Sample Collection Activities

- Date(s) and times of the sample collection or event.
- Number and types of collected samples.
- Sample location with an emphasis on any changes to documentation in governing documents (i.e., SAP, FSP). This may include measurements from reference points or sketches of sample locations with respect to local features.
- Sample identification numbers, including any applicable cross-references to split samples or samples collected by another entity.
- A description of sampling methodology, or reference to any governing document (i.e., FSP, SAP, QAPP).
- Summary of equipment preparation and decontamination procedures.
- Sample description including depth, color, texture, moisture content, and evidence of waste material or staining.
- Air monitoring (field screening) results.
- Types of laboratory analyses requested.

Site Health and Safety Activities

• All safety, accident, and/or incident reports.

SOP	1501.01				
GROUP	Field Documentation	on			
SUB-GROUP					
TITLE	Field Logbook				
DATE	11/19/2001	FILE	1501-01.DOC	PAGE	3 of 3

- Real-time personnel air monitoring results, if applicable, or if not documented in the HASP.
- Heat/cold stress monitoring data, if applicable.
- Reasons for upgrades or downgrades in personal protective equipment.
- Health and safety inspections, checklists (drilling safety guide), meetings/briefings.
- Calibration records for field instruments.

Oversight Activities

- Progress and activities performed by contractors including operating times.
- Deviations of contractor activities with respect to project governing documents (i.e., specifications).
- Contractor sampling results and disposition of contingent soil materials/stockpiles.
- Excavation specifications and locations of contractor confirmation samples.
- General site housekeeping and safety issues by site contractors.

SOP	1502.01				
GROUP	Field Documentation	on			
SUB-GROUP					
TITLE	Photograph Logs				
DATE	11/19/2001	FILE	1502-01.DOC	PAGE	1 of 1

The following Standard Operating Procedure (SOP) presents the requirements for collecting information related to photodocumentation of site activities.

PROCEDURE

- Uniquely number each roll of film obtained for use.
- Record the following information for each negative exposed:
 - 1. Date and Time
 - 2. Photographer Name
 - 3. Witness Name
 - 4. Orientation (Landscape, Portrait, or Panaoramic)
 - 5. Description (including activity being performed, specific equipment of interest, sample location(s), compass direction photographer is facing)
- Record "NA" for the negatives not used if the roll is not completely used prior to development.
- Record unique roll number on receipt when film is submitted for development.
- Verify descriptions on log with negative numbers when photographs are received from processing.

FORMS

Blank Photograph Logs can be printed from WESTON On-Line from the *Records Management Application*. Selecting the *Reports/Project Planning/Blank Photo Logs* menu option will generate a project specific log with 36 entries.

SOP	0110.01					
GROUP	Database Management System					
SUB-GROUP	Data Collection a	Data Collection and Acquisition				
TITLE	Sample Nomenclature					
DATE	02/26/2009	FILE	0110-20060227.DOC	PAGE	1 of 2	

The following Standard Operating Procedure (SOP) presents the sample nomenclature for analytical samples that will generate unique sample names compatible with most data management systems. The sample nomenclature is based upon specific requirements for the reporting of these results. A site specific data management plan should be prepared prior to sample collection.

PROCEDURE

SAMPLE NOMENCLATURE - SOIL AND SEDIMENT

Area of Concern – ID – Depth - Collection Type + QC Type

Where:

Area of Concern: A four-digit identifier used to designate the particular Area of Concern

(AOC) that the location where the sample was collected.

ID: A three-digit identifier used to designate the particular location in the AOC

from which the sample was collected or the center of the composite sample.

Depth: A two-digit code used to designate what depth of sample was collected:

03	0 to 3 inches
06	3 to 6 inches
12	6 to 12 inches

Collection Type: A one-digit code used to designate what type of sample was collected:

1	Surface Water
2	Ground Water
3	Leachate
4	Field QC/water sample
5	Soil/Sediment

6	Oil
7	Waste
8	Other
9	Drinking Water

QC Type: A one-digit code used to designate the QC type of the sample:

1	Normal
2	Duplicate
3	Rinsate Blank
4	Trip Blank
5	Field Blank
6	Confirmation

Examples:

- 2054-055-06-51: Represents the normal soil sample collected from AOC 2054 at location 055 from 3 to 6 inches of depth.
- **2054-055-06-52**: Represents the duplicate soil sample collected from AOC 2054 at location 055 from 3 to 6 inches of depth.
- 2054-055-06-43: Represents the rinsate water sample collected after the last sample of the day if last sample was collected from AOC 2054 at location 055 from 3 to 6 inches of depth.

SOP	0110.01					
GROUP	Database Management System					
SUB-GROUP	Data Collection a	Data Collection and Acquisition				
TITLE	Sample Nomenclature					
DATE	02/26/2009	FILE	0110-20060227.DOC	PAGE	2 of 2	

SAMPLE NOMENCLATURE – WATER (from fixed station or location to be sampled more than once)

WELL OR STATION – YYYYMMDD - Collection Type + QC Type

Where:

Well or Station: For Wells and boreholes always assume there will be 10 or more so

Monitoring Well 1 becomes designated MW01 or MW-01. If it is anticipated that there will be over 100 wells designate Monitoring Well 1 as MW001 or

MW-001.

YYYYMMDD: A four-digit year + two-digit month + two-digit day

Collection Type: A one-digit code used to designate what type of sample was collected and are

shown on page 1.

QC Type: A one-digit code used to designate the QC type of the sample and are shown

on page 1.

Examples:

• *MW01-20090226-21:* Represents the normal groundwater sample collected from Monitoring Well 1 on 26 February 2009.

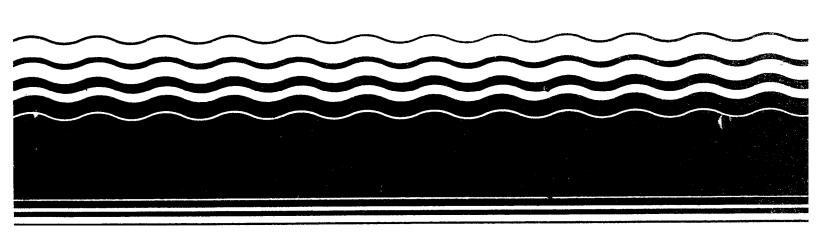
• *MW01-20090226-44:* Represents the trip blank in the same ice chest as the groundwater sample in the previous collected from Monitor Well 1 on 02/26/2009. All trip blanks must have a sample ID and they must be unique and on the Chain-of -Custody.

• 2054-000-00-43: Represents the rinsate sample from AOC 2054

United States Environmental Protection Agency Office of Solid Waste and Emergency Response Washington DC 20460 PB91-921275 EPA/540/P-91/007 January 1999 OSWER 9360.4-06

SEPA

Compendium of ERT Groundwater Sampling Procedures



COMPENDIUM OF ERT GROUNDWATER SAMPLING PROCEDURES

Sampling Equipment Decontamination

Groundwater Well Sampling

Soil Gas Sampling

Monitoring Well Installation

Water Level Measurement

Well Development

Controlled Pumping Test

Slug Test

Interim Final

Environmental Response Team Emergency Response Division

Office of Emergency and Remedial Response
U.S. Environmental Protection Agency
Washington, DC 20460



Notice

This document has been reviewed in accordance with U.S. Environmental Protection Agency policy and approved for publication. Mention of trade names or commercial products does not constitute endorsement or recommendation for use.

The policies and procedures established in this document are intended solely for the guidance of government personnel, for use in the Superfund Removal Program. They are not intended, and cannot be relied upon, to create any rights, substantive or procedural, enforceable by any party in litigation with the United States. The Agency reserves the right to act at variance with these policies and procedures and to change them at any time without public notice.

Depending on circumstances and needs, it may not be possible or appropriate to follow these procedures exactly in all situations due to site conditions, equipment limitations, and limitations of the standard procedures. Whenever these procedures cannot be followed as written, they may be used as general guidance with any and all modifications fully documented in either QA Plans, Sampling Plans, or final reports of results.

Each Standard Operating Procedure in this compendium contains a discussion on quality assurance/quality control (QA/QC). For more information on QA/QC objectives and requirements, refer to the *Quality Assurance/Quality Control Guidance for Removal Activities*, OSWER directive 9360.4-01, EPA/540/G-90/004.

Questions, comments, and recommendations are welcomed regarding the Compendium of ERT Groundwater Sampling Procedures. Send remarks to:

Mr. William A. Coakley Removal Program QA Coordinator U.S. EPA - ERT Raritan Depot - Building 18, MS-101 2890 Woodbridge Avenue Edison, NJ 08837-3679

For additional copies of the Compendium of ERT Groundwater Sampling Procedures, please contact:

National Technical Information Service (NTIS)
U.S. Department of Commerce
5285 Port Royal Road
Springfield, VA 22161
(703) 487-4600

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Acknowledgments

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1.0 SAMPLING EQUIPMENT DECONTAMINATION: SOP #2006

1.1 SCOPE AND APPLICATION

This Standard Operating Procedure (SOP) describes methods used for preventing or reducing cross-contamination, and provides general guidelines for sampling equipment decontamination procedures at a hazardous waste site. Preventing or minimizing cross-contamination in sampled media and in samples is important for preventing the introduction of error into sampling results and for protecting the health and safety of site personnel.

Removing or neutralizing contaminants that have accumulated on sampling equipment ensures protection of personnel from permeating substances, reduces or eliminates transfer of contaminants to clean areas, prevents the mixing of incompatible substances, and minimizes the likelihood of sample cross-contamination.

1.2 METHOD SUMMARY

Contaminants can be physically removed from equipment, or deactivated by sterilization or disinfection. Gross contamination of equipment requires physical decontamination, including abrasive and non-abrasive methods. These include the use of brushes, air and wet blasting, and high-pressure water cleaning, followed by a wash/rinse process using appropriate cleaning solutions. Use of a solvent rinse is required when organic contamination is present.

1.3 SAMPLE PRESERVATION, CONTAINERS, HANDLING, AND STORAGE

This section is not applicable to this SOP.

1.4 INTERFERENCES AND POTENTIAL PROBLEMS

The use of distilled/deionized water commonly available from commercial vendors may be acceptable for decontamination of sampling equipment

provided that it has been verified by laboratory analysis to be analyte free.

- An untreated potable water supply is not an acceptable substitute for tap water. Tap water may be used from any municipal water treatment system for mixing of decontamination solutions.
- Acids and solvents utilized in the decontamination sequence pose the health and safety risks of inhalation or skin contact, and raise shipping concerns of permeation or degradation.
- The site work plan must address disposal of the spent decontamination solutions.
- Several procedures can be established to minimize contact with waste and the potential for contamination. For example:

Stress work practices that minimize contact with hazardous substances.

Use remote sampling, handling, and container-opening techniques when appropriate.

Cover monitoring and sampling equipment with protective material to minimize contamination.

Use disposable outer garments and disposable sampling equipment when appropriate.

1.5 EQUIPMENT/APPARATUS

- appropriate personal protective clothing
- non-phosphate detergent
- selected solvents
- long-handled brushes
- drop cloths/plastic sheeting
- trash container
- paper towels
- galvanized tubs or buckets
- tap water

- distilled/deionized water
- metal/plastic containers for storage and disposal of contaminated wash solutions
- pressurized sprayers for tap and deionized/distilled water
- sprayers for solvents
- trash bags
- aluminum foil
- safety glasses or splash shield
- emergency eyewash bottle

1.6 REAGENTS

There are no reagents used in this procedure aside from the actual decontamination solutions and solvents. In general, the following solvents are utilized for decontamination purposes:

- zz 10% nitric acid(1)
- zz acetone (pesticide grade)(*)
- ** hexane (pesticide grade)(*)
- (1) Only if sample is to be analyzed for trace metals.
- (2) Only if sample is to be analyzed for organics.

1.7 PROCEDURES

As part of the health and safety plan, develop and set up a decontamination plan before any personnel or equipment enter the areas of potential exposure. The equipment decontamination plan should include:

- the number, location, and layout of decontamination stations
- which decontamination apparatus is needed
- the appropriate decontamination methods
- methods for disposal of contaminated clothing, apparatus, and solutions

1.7.1 Decontamination Methods

All personnel, samples, and equipment leaving the contaminated area of a silt must be decontaminated. Various decontamination methods will either physically remove contaminants, inactivate contaminants by disinfection or sterilization, or do hot h.

In many cases, gross contamination can be removed by physical means. The physical decontamination techniques appropriate for equipment decontamination can be grouped into two categories: abrasive methods and non-abrasive methods.

Abrasive Cleaning Methods

Abrasive cleaning methods work by rubbing and wearing away the top layer of the surface containing the contaminant. The following abrasive methods are available:

- Mechanical: Mechanical cleaning methods use brushes of metal or nylon. The amount and type of contaminants removed will vary with the hardness of bristles, length of brushing time, and degree of brush contact.
- Air Blasting: Air blasting is used for cleaning large equipment, such as bulldozers, drilling rigs or auger bits. The equipment used in air blast cleaning employs compressed air to force abrasive material through a nozzle at high velocities. The distance between the nozzle and the surface cleaned, as well as the pressure of air, the time of application, and the angle at which the abrasive strikes the surface, determines cleaning efficiency. Air blasting has several disadvantages: it is unable to control the amount of material removed, it can acrate contaminants, and it generates large amounts of waste.
- Wet Blasting: Wet blast cleaning, also used to clean large equipment, involves use of a suspended fine abrasive: delivered by compressed air to the contaminated area. The amount of materials removed can be carefully controlled by using wry line abrasives. This method generates a large amount of waste.

Non-Abrasive Cleaning Methods

Non-abrasive cleaning methods work by forcing the contaminant off of a surface with pressure. In general, less of the equipment surface is removed using non-ahrasive methods. The following non-abrasive methods are available:

- High-Pressure Water: This method consists of a high-pressure pump, an operator-controlled directional nozzle, and a high pressure hose. Operating pressure usually ranges from 340 to 680 atmospheres (atm) which relates to flow rates of 20 to 140 liters per minute.
- Ultra-High-Pressure Water: This system produces a pressurized water jet (from 1,000 to 4,000 atm). The ultra-high-pressure spray removes tightly-adhered surface film. The water velocity ranges from 500 m/sec (1,000 atm) to 900 m/sec (4,000 atm). Additives can enhance the method. This method is not applicable for hand-held sampling equipment.

Disinfection/Rinse Methods

- Disinfection: Disinfectants are a practical means of inactivating infectious agents.
- Sterilization: Standard sterilization methods involve heating the equipment.

 Sterilization is impractical for large equipment.
- Rinsing: Rinsing removes contaminants through dilution, physical attraction, and solubilization.

1.7.2 Field Sampling Equipment Cleaning Procedures

Solvent rinses are not necessarily required when organics are not a contaminant of concern and may be eliminated from the sequence specified below. Similarly, an acid rinse is not required if analysis does not include inorganics.

- 1. Where applicable, follow physical removal procedures specified in section 1.7.1.
- 2. Wash equipment with a non-phosphate detergent solution.
- 3. Rinse with tap water.
- 4. Rinse with distilled/deionized water.
- 5. Rinse with 10% nitric acid if the sample will be analyzed for trace organics.

- 6. Rinse with distilled/deionized water.
- 7. Use a solvent rinse (pesticide grade) if the sample will be analyzed for organics.
- 8. Air dry the equipment completely.
- 9. Rinse again with distilled/deionized water.

Selection of the solvent for use in the decontamination process is based on the contaminants present at the site. Use of a solvent is required when organic contamination is present on-site. Typical solvents used for removal of organic contaminants include acetone, hexane, or water. An acid rinse step is required if metals are present on-site. If a particular contaminant fraction is not present at the site, the nine-step decontamination procedure listed above may be modified for site specificity. The decontamination solvent used should not be among the contaminants of concern at the site.

Table 1 lists solvent rinses which may be required for elimination of particular chemicals. After each solvent rinse, the equipment should be air dried and rinsed with distilled/deionized water.

Sampling equipment that requires the use of plastic tubing should be disassembled and the tubing replaced with clean tubing, before commencement of sampling and between sampling locations.

1.8 CALCULATIONS

This section is not applicable to this SOP.

1.9 QUALITY ASSURANCE/ QUALITY CONTROL

One type of quality control sample specific to the field decontamination process is the rinsate blank. The rinsate blank provides information on the effectiveness of the decontamination process employed in the field. When used in conjunction with field blanks and trip blanks, a rinsate blank can detect contamination during sample handling, storage and sample transportation to the laboratory.

Table 1: Recommended Solvent Rinse for Soluble Contaminants

SOLVENT	SOLUBLE CONTAMINANTS
Water	Low-chain hydrocarbons Inorganic compounds Salts Some organic acids and other polar compounds
Dilute Acids	Basic (caustic) compounds Amines Hydrazines
Dilute Bases for example, detergent and soap	 Metals Acidic compounds Phenol Thiols Some nitro and sulfonic compounds
Organic Solvents(1) - for example, alcohols, ethers, ketones, aromatics, straight-chain alkanes (e.g., hexane), and common petroleum products (e.g., fuel, oil, kerosene)	Nonpolar compounds (e.g., some organic compounds)

(1) - WARNING: Some organic solvents can permeate and/or degrade protective clothing.

A rinsate blank consists of a sample of analyte-free (i.e, deionized) water which is passed over and through a field decontaminated sampling device and placed in a clean sample container.

Rinsate blanks should be run for all parameters of interest at a rate of 1 per 20 for each parameter, even if samples are not shipped that day. Rinsate blanks are not required if dedicated sampling equipment is used.

1.10 DATA VALIDATION

This section is not applicable to this SOP.

1.11 HEALTH AND SAFETY

When working with potentially hazardous materials, follow U.S. EPA, OSHA and specific health and safety procedures.

Decontamination can pose hazards under certain circumstances even though performed to protect

health and safety. Hazardous substances may be incompatible with decontamination methods. For example, the decontamination solution or solvent may react with contaminants to produce heat, explosion, or toxic products. Decontamination methods may be incompatible with clothing or equipment; some solvents can permeate or degrade protective clothing. Also, decontamination solutions and solvents may pose a direct health hazard to workers through inhalation or skin contact, or if they combust.

The decontamination solutions and solvents must be determined to be compatible before use. Any method that permeates, degrades, or damages personal protective equipment should not be used. If decontamination methods pose a direct health hazard, measures should be taken to protect personnel or the methods should be modified to eliminate the hazard.

2.0 GROUNDWATER WELL SAMPLING: SOP #2007

2.1 SCOPE AND APPLICATION

The objective of this Standard Operating Procedure (SOP) is to provide general reference information on sampling of groundwater wells. This guideline is primarily concerned with the collection of water samples from the saturated zone of the subsurface. Every effort must be made to ensure that the sample is representative of the particular zone of water being sampled. These procedures are designed to be used in conjunction with analyses for the most common types of groundwater contaminants (e.g., volatile and semi-volatile organic compounds, pesticides, metals, biological parameters).

2.2 METHOD SUMMARY

Prior to sampling a monitoring well, the well must be purged. This may be done with a number of instruments. The most common of these are the bailer, submersible pump, non-gas contact bladder pump and inertia pump. At a minimum, three well volumes should be purged, if possible. Equipment must be decontaminated prior to use and between wells. Once purging is completed and the correct laboratory-cleaned sample containers have been prepared, sampling may proceed. Sampling may be conducted with any of the above instruments, and need not be the same as the device used for purging. Care should be taken when choosing the sampling device as some will affect the integrity of the sample. Sampling equipment must also be Sampling should occur in a decontaminated. progression from the least to most contaminated well, if this information is known.

2.3 SAMPLE PRESERVATION, CONTAINERS, HANDLING, AND STORAGE

The type of analysis for which a sample is being collected determines the type of bottle, preservative, holding time, and filtering requirements. Samples should be collected directly from the sampling device into appropriate laboratory-cleaned containers. Check that a Teflon liner is present in

the cap, if required. Attach a sample identification label. Complete a field data sheet, a chain of custody form and record all pertinent data in the site logbook.

Samples shall be appropriately preserved, labelled, logged, and placed in a cooler to be maintained at 4°C. Samples must be shipped well before the holding time is over and ideally should be shipped within 24 hours of sample collection. It is imperative that these samples be shipped or delivered daily to the analytical laboratory in order to maximize the time available for the laboratory to perform the analysis. The bottles should be shipped with adequate packing and cooling to ensure that they arrive intact.

Certain conditions may require special handling techniques. For example, treatment of a sample for volatile organic (WA) analysis with sodium thiosulfate preservative is required if there is residual chlorine in the water (such as public water supply) that could cause free radical chlorination and change the identity of the original contaminants. However, sodium thiosulfate should not be used if chlorine is not present in the water. Special requirements must be determined prior to conducting fieldwork.

2.4 INTERFERENCES AND POTENTIAL PROBLEMS

2.4.1 General

The primary goal of groundwater sampling is to obtain a representative sample of the groundwater body. Analysis can be compromised by field personnel in two primary ways: (1) taking an unrepresentative sample, or (2) by incorrect handling of the sample. There are numerous ways of introducing foreign contaminants into a sample, and these must be avoided by following strict sampling procedures and only utilizing trained field personnel.

2.4.2 Purging

In a non-pumping well, there will be little or no vertical mixing of the water, and stratification will

occur. The well water in the screened section will mix with the groundwater due to normal flow patterns, but the well water above the screened section will remain isolated, become stagnant and lack the VOAs representative of the groundwater. Sampling personnel should realize that stagnant water may contain foreign material inadvertently or deliberately introduced from the surface, resulting in an unrepresentative sample. To safeguard against collecting nonrepresentative stagnant water, follow these guidelines during sampling:

- As a general rule, all monitoring wells should be pumped or bailed prior to Purge water should be sampling. containerized on site or handled as specified in the site-specific project plan. Evacuation of a minimum of one volume of water in the well casing, and preferably three to five volumes, is recommended for a representative sample. In a high-yielding ground water formation and where there is no stagnant water in the well above the screened section, evacuation prior to sample withdrawal is not as critical. However, in all cases where the monitoring data is to be used for enforcement actions, evacuation is recommended.
- For wells that can be pumped or bailed to dryness with the equipment being used, the well should be evacuated and allowed to recover prior to sample withdrawal. If the recovery rate is fairly rapid and the schedule allows, evacuation of more than one volume of water is preferred. If recovery is slow, sample the well upon recovery after one evacuation.
- A nonrepresentative sample can also result from excessive pre-pumping of the monitoring well. Stratification of the leachate concentration in the groundwater formation may occur, or heavier-than-water compounds may sink to the lower portions of the aquifer. Excessive pumping can dilute or increase the contaminant concentrations from what is representative of the sampling point of interest.

2.4.3 Materials

Samplers and evacuation equipment (bladders, pumps, bailers, tubing, etc.) should be limited to

those made with stainless steel, Teflon, and glass in areas where concentrations are expected to be at or near the detection limit. The tendency of organics to leach into and out of many materials make the selection of materials critical for trace analyses. The use of plastics, such as PVC or polyethylene, should be avoided when analyzing for organics. However, PVC may be used for evacuation equipment as it will not come in contact with the sample.

Table 2 on page 7 discusses the advantages and disadvantages of certain equipment.

2.5 EQUIPMENT/APPARATUS

2.5.1 General

- water level indicator
 - electric sounder
 - steel tape
 - transducer
 - reflection sounder
 - airline
- depth sounder
- appropriate keys for well cap locks
- steel brush
- HNU or OVA (whichever is most appropriate)
- logbook
- calculator
- field data sheets
- chain of custody forms
- forms and seals
- sample containers
- Engineer's rule
- sharp knife (locking blade)
- tool box (to include at least: screwdrivers, pliers, hacksaw, hammer, flashlight, adjustable wrench)
- leather work gloves
- appropriate health and safety gear
- 5-gallon pail
- plastic sheeting
- shipping containers
- packing materials
- bolt cutters
- Ziploc plastic bags
- · containers for evacuation of liquids
- decontamination solutions
- tap water
- non-phosphate soap
- several brushes

Table 2: Advantages and Disadvantages of Various Groundwater Sampling Devices

Device	Advantages	Disadvantages
Bailer	The only practical limitations are size and materials No power source needed Portable Inexpensive; it can be dedicated and hung in a well reducing the chances of crosscontamination Minimal outgassing of volatile organics while sample is in bailer Readily available Removes stagnant water first Rapid, simple method for removing small volumes of purge water	Time consuming, especially for large wells Transfer of sample may cause aeration
Submersible Pump	Portable; can be used on an unlimited number of wells Relatively high pumping rate (dependent on depth and size of pump) Generally very reliable; does not require priming	Potential for effects on analysis of trace organics Heavy and cumbersome, particularly in deeper wells Expensive Susceptible to damage from silt or sediment Impractical in low yielding or shallow wells
Non-Gas Contact Bladder Pump	Maintains integrity of sample Easy to use	Difficult to clean although dedicated tubing and bladder may be used Only useful to approximately 100 feet in depth Supply of gas for operation (bottled gas and/or compressor) is difficult to obtain and is cumbersome
Suction Pump	EX Portable, inexpensive, and readily available	Only useful to approximately 25 feet or less in depth Vacuum can cause loss of dissolved gases and volatile organics Pump must be primed and vacuum is often difficult to maintain May cause pH modification
Inertia Pump	Portable, inexpensive, and readily available Rapid method for purging relatively shallow wells	Only useful to approximately 70 feet or less in depth May be time consuming to use Labor intensive WaTerra pump is only effective in 2-inch diameter wells

- pails or tubs
- aluminum foil
- garden sprayer
- preservatives
- distilled or deionized water

2.5.2 **Bailer**

- clean, decontaminated bailer(s) of appropriate size and construction material
- nylon line, enough to dedicate to each well
- Teflon-coated bailer wire
- sharp knife
- aluminum foil (to wrap clean bailers)
- 5-gallon bucket

2.5.3 Submersible Pump

- pump(s)
- generator (110, 120, or 240 volt) or 12-volt battery if inaccessible to field vehicle
- l-inch black PVC coil pipe -- enough to dedicate to each well
- hose clamps
- safety cable
- tool box supplement
 - pipe wrenches, 2
 - wire strippers
 - electrical tape
 - heat shrink
 - hose connectors
 - Teflon tape
- winch or pulley
- gasoline for generator
- flow meter with gate valve
- l-inch nipples and various plumbing (i.e., pipe connectors)

2.5.4 Non-Gas Contact Bladder Pump

- non-gas contact bladder pump
- compressor or nitrogen gas tank
- batteries and charger
- Teflon tubing -- enough to dedicate to each well
- Swagelock fitting
- toolbox supplements -- same as submersible pump

2.5.5 Suction Pump

- pump
- black coil tubing -- enough to dedicate to each well

- gasoline -- if required
- toolbox
- zz plumbing fittings

2.5.6 Inertia Pump

- pump assembly (WaTerra pump, piston Pump)
- ≤ 5-gallon bucket

2.6 REAGENTS

Reagents will be utilized for preservation of samples and for decontamination of sampling equipment. The preservation required is specified by the analysis to be performed. Decontamination solutions are specified in ERT SOP #2006, Sampling Equipment Decontamination.

2.7 PROCEDURES

2.7.1 Preparation

- 1. Determine the extent of the sampling effort, the sampling methods to be employed, and which equipment and supplies are needed.
- 2. Obtain necessary sampling and monitoring equipment.
- 3. Decontaminate or preclean equipment, and ensure that it is in working order.
- 4. Prepare scheduling and coordinate with staff, clients, and regulatory agency, if appropriate.
- Perform a general site survey prior to site entry in accordance with the site-specific health and safety plan.
- 6. Identify and mark all sampling locations.

2.7.2 Field Preparation

- 1. Start at the least contaminated well, if known.
- 2. Lay plastic sheeting around the well to minimize likelihood of contamination of equipment from soil adjacent to the well.

- 3. Remove locking well cap, note location, time of day, and date in field notebook or an appropriate log form.
- 4. Remove well casing cap.
- Screen headspace of well with an appropriate monitoring instrument to determine the presence of volatile organic compounds and record in site logbook.
- 6. Lower water level measuring device or equivalent (i.e., permanently installed transducers or airline) into well until water surface is encountered.
- 7. Measure distance from water surface to reference measuring point on well casing or protective barrier post and record in site logbook. Alternatively, if there is no reference point, note that water level measurement is from top of steel casing, top of PVC riser pipe, from ground surface, or some other position on the well head.
- 8. Measure total depth of well (do this at least twice to confirm measurement) and record in site logbook or on log form.
- 9. Calculate the volume of water in the well and the volume to be purged using the calculations in Section 2.8.
- 10. Select the appropriate purging and sampling equipment.

2.7.3 Evacuation of Static Water (Purging)

The amount of flushing a well receives prior to sample collection depends on the intent of the monitoring program as well as the hydrogeologic conditions. Programs where overall quality determination of water resources are involved may require long pumping periods to obtain a sample that is representative of a large volume of that aquifer. The pumped volume can be determined prior to sampling so that the sample is a composite of known volume of the aquifer, or the well can be pumped until the stabilization of parameters such as temperature, electrical conductance, or pH has occurred.

However, monitoring for defining a contaminant plume requires a representative sample of a small volume of the aquifer. These circumstances require that the well be pumped enough to remove the stagnant water but not enough to induce flow from other areas. Generally, three well volumes are considered effective, or calculations can be made to determine, on the basis of the aquifer parameters and well dimensions, the appropriate volume to remove prior to sampling.

During purging, water level measurements may be taken regularly at 15- to 30-second intervals. This data may be used to compute aquifer transmissivity and other hydraulic characteristics.

The following well evacuation devices are most commonly used. Other evacuation devices are available, but have been omitted in this discussion due to their limited use.

Bailer

Bailers are the simplest purging device used and have many advantages. They generally consist of a rigid length of tube, usually with a ball check-valve at the bottom. A line is used to lower the bailer into the well and retrieve a volume of water. The three most common types of bailer are PVC, Teflon, and stainless steel.

This manual method of purging is best suited to shallow or narrow diameter wells. For deep, larger diameter wells which require evacuation of large volumes of water, other mechanical devices may be more appropriate.

Bailing equipment includes a clean decontaminated bailer, Teflon or nylon line, a sharp knife, and plastic sheeting.

- 1. Determine the volume of water to be purged as described in Section 2.7.2, Field Preparation.
- Lay plastic sheeting around the well to prevent contamination of the bailer line with foreign materials.
- 3. Attach the line to the bailer and lower until the bailer is completely submerged.
- 4. Pull bailer out ensuring that the line either falls onto a clean area of plastic sheeting or never touches the ground.

- 5. Empty the bailer into a pail until full to determine the number of bails necessary to achieve the required purge volume.
- Thereafter, pour the water into a container and dispose of purge waters as specified in the sitespecific project plan.

Submersible Pump

Submersible pumps are generally constructed of plastic, rubber, and metal parts which may affect the analysis of samples for certain trace organics and inorganics. As a consequence, submersible pumps may not be appropriate for investigations requiring analyses of samples for trace contaminants. However, they are still useful for pre-sample purging. However, the pump must have a check valve to prevent water in the pump and the pipe from rushing back into the well.

Submersible pumps generally use one of two types of power supplies, either electric or compressed gas. Electric pumps can be powered by a 12-volt DC rechargeable battery, or a 110- or 220-volt AC power supply. Those units powered by compressed gas normally use a small electric compressor which also needs 12-volt DC or 110-volt AC power. They may also utilize compressed gas from bottles. Pumps differ according to the depth and diameter of the monitoring wells.

- 1. Determine the volume of water to be purged as described in section 2.7.2, Field Preparation.
- Lay plastic sheeting around the well to prevent contamination of pumps, hoses or lines with foreign materials.
- 3. Assemble pump, hoses and safety cable, and lower the pump into the well. Make sure the pump is deep enough so that purging does not evacuate all the water. (Running the pump without water may cause damage.)
- 4. Attach flow meter to the outlet hose to measure the volume of water purged.
- 5. Attach power supply, and purge well until specified volume of water has been evacuated (or until field parameters, such as temperature, pH, conductivity, etc. have stabilized). Do not allow the pump to run dry. If the pumping rate

- exceeds the well recharge rate, lower the pump further into the well, and continue pumping.
- 6. Collect and dispose of purge waters as specified in the site-specific project plan.

Non-Contact Gas Bladder Pump

For this procedure, an all stainless-steel and Teflon Middleburg-squeeze bladder pump (e.g., IEA, TIMCO, Well Wizard, Geoguard, and others) is used to provide the least amount of material interference to the sample (Barcelona, 1985). Water comes into contact with the inside of the bladder (Teflon) and the sample tubing, also Teflon, that may be dedicated to each well. Some wells may have permanently installed bladder pumps (i.e., Well Wizard, Geoguard), that will be used to sample for all parameters.

- Assemble Teflon tubing, pump and charged control box.
- 2. Use the same procedure for purging with a bladder pump as for a submersible pump.
- 3. Be sure to adjust flow rate to prevent violent jolting of the hose as sample is drawn in.

Suction Pump

There are many different types of suction pumps. They include: centrifugal, peristaltic and diaphragm. Diaphragm pumps can be used for well evacuation at a fast pumping rate and sampling at a low pumping rate. The peristaltic pump is a low-volume pump that uses rollers to squeeze the flexible tubing, thereby creating suction. This tubing can be dedicated to a well to prevent cross-contamination. Peristaltic pumps, however, require a power source.

- 1. Assemble the pump, tubing, and power source if necessary.
- 2. To purge with a suction pump, follow the exact procedures outlined for the submersible pump.

Inertia Pump

Inertia pumps, such as the WaTerra pump and piston pump, are manually operated. They are appropriate to use when wells are too deep to bail by hand, but are not inaccessible enough to warrant an automatic (submersible, etc.) pump. These

pumps are made of plastic and may be either decontaminated or discarded, after use.

- 1. Determine the volume of water to be purged as described in Section 2.7.2, Field Preparation.
- Lay plastic sheeting around the well to prevent contamination of pumps or hoses with foreign materials.
- 3. Assemble pump, and lower to the appropriate depth in the well.
- Begin pumping manually, discharging water into a 5-gallon bucket (or other graduated vessel). Purge until specified volume of water has been evacuated (or until field parameters such as temperature, pH, conductivity, etc. have stabilized).
- 5. Collect and dispose of purge waters as specified in the site-specific project plan.

2.7.4 Sampling

Sample withdrawal methods require the use of pumps, compressed air, bailers, and samplers. Ideally, purging and sample withdrawal equipment should be completely inert, economical to use, easily cleaned, sterilized, reusable, able to operate at remote sites in the absence of power resources, and capable of delivering variable rates for sample collection.

There are several factors to take into consideration when choosing a sampling device. Care should be taken when reviewing the advantages or disadvantages of any one device. It may be appropriate to use a different device to sample than that which was used to purge. The most common example of this is the use of a submersible pump to purge and a bailer to sample.

Bailer

The positive-displacement volatile sampling bailer (by GPI) is perhaps the most appropriate for collection of water samples for volatile analysis. Other bailer types (messenger, bottom fill, etc.) are less desirable, but may be mandated by cost and site conditions. Generally, bailers can provide an acceptable sample, providing that sampling personnel use extra care in the collection process.

- Surround the monitoring well with clean plastic sheeting.
- 2. Attach a line to the bailer. If a bailer was used for purging, the same bailer and line may be used for sampling.
- 3. Lower the bailer slowly and gently into the well, taking care not to shake the casing sides or to splash the bailer into the water. Stop lowering at a point adjacent to the screen.
- 4. Allow bailer to fill and then slowly and gently retrieve the bailer from the well, avoiding contact with the casing, so as not to knock flakes of rust or other foreign materials into the bailer.
- Remove the cap from the sample container and place it on the plastic sheet or in a location where it will not become contaminated. See Section 2.7.7 for special considerations on VOA samples.
- 6. Begin pouring slowly from the bailer.
- 7. Filter and preserve samples as required by sampling plan.
- 8. Cap the sample container tightly and place prelabeled sample container in a carrier.
- 9. Replace the well cap.
- 10. Log all samples in the site logbook and on field data sheets and label all samples.
- 11. Package samples and complete necessary paperwork.
- 12. Transport sample to decontamination zone to prepare it for transport to analytical laboratory.

Submersible Pump

Although it is recommended that samples not be collected with a submersible pump due to the reasons stated in Section 2.4, there are some situations where they may be used.

1. Allow the monitoring well to recharge after purging, keeping the pump just above the screened section.

- 2. Attach gate valve to hose (if not already fitted), and reduce flow of water to a manageable sampling rate.
- 3. Assemble the appropriate bottles.
- 4. If no gate valve is available, run the water down the side of a clean jar and fill the sample bottles from the jar.
- 5. Cap the sample container tightly and place prelabeled sample container in a carrier.
- 6. Replace the well cap.
- 7. Log all samples in the site logbook and on the field data sheets and label all samples.
- 8. Package samples and complete necessary paperwork.
- Transport sample to decontamination zone for preparation for transport to analytical laboratory.
- 10. Upon completion, remove pump and assembly and fully decontaminate prior to setting into the next sample well. Dedicate the tubing to the hole.

Non-Gas Contact Bladder Pump

The use of a non-gas contact positive displacement bladder pump is often mandated by the use of dedicated pumps installed in wells. These pumps are also suitable for shallow (less than 100 feet) wells. They are somewhat difficult to clean, but may be used with dedicated sample tubing to avoid cleaning. These pumps require a power supply and a compressed gas supply (or compressor). They may be operated at variable flow and pressure rates making them ideal for both purging and sampling.

Barcelona (1984) and Nielsen (1985) report that the non-gas contact positive displacement pumps cause the least amount of alteration in sample integrity as compared to other sample retrieval methods.

- 1. Allow well to recharge after purging.
- 2. Assemble the appropriate bottles.

- 3. Turn pump on, increase the cycle time and reduce the pressure to the minimum that will allow the sample to come to the surface.
- 4. Cap the sample container tightly and place prelabeled sample container in a carrier.
- 5. Replace the well cap.
- Log all samples in the site logbook and on field data sheets and label all samples.
- 7. Package samples and complete necessary paperwork.
- Transport sample to decontamination zone for preparation for transport to analytical laboratory.
- On completion, remove the tubing from the well and either replace the Teflon tubing and bladder with new dedicated tubing and bladder or rigorously decontaminate the existing materials.
- 10. Collect non-filtered samples directly from the outlet tubing into the sample bottle.
- 11. For filtered samples, connect the pump outlet tubing directly to the filter unit. The pump pressure should remain decreased so that the pressure build-up on the filter does not blow out the pump bladder or displace the filter. For the Geotech barrel filter, no actual connections are necessary so this is not a concern.

Suction Pump

In view of the limitations of suction pumps, they are not recommended for sampling purposes.

Inertia Pump

Inertia pumps may be used to collect samples. It is more common, however, to purge with these pumps and sample with a bailer.

- 1. Following well evacuation, allow the well to recharge.
- 2. Assemble the appropriate bottles.

- 3. Since these pumps are manually operated, the flow rate may be regulated by the sampler. The sample may be discharged from the pump outlet directly into the appropriate sample container.
- 4. Cap the sample container tightly and place prelabeled sample container in a carrier.
- 5. Replace the well cap.
- 6. Log all samples in the site logbook and on field data sheets and label all samples.
- 7. Package samples and complete necessary paperwork.
- 8. Transport sample to decontamination zone for preparation for transport to analytical laboratory.
- 9. Upon completion, remove pump and decontaminate or discard, as appropriate.

2.7.5 Filtering

For samples that require filtering, such as samples which will be analyzed for total metals, the filter must be decontaminated prior to use and between uses. Filters work by two methods. A barrel filter such as the "Geotech" filter works with a bicycle pump, which is used to build up positive pressure in the chamber containing the sample. The sample is then forced through the filter paper (minimum size 0.45μ) into a jar placed underneath. The barrel itself is filled manually from the bailer or directly via the hose of the sampling pump. The pressure must be maintained up to 30 psi by periodic pumping.

A vacuum type filter involves two chambers, the upper chamber contains the sample and a filter (minimum size 0.45μ) divides the chambers. Using a hand pump or a Gilian type pump, air is withdrawn from the lower chamber, creating a vacuum and thus causing the sample to move through the filter into the lower chamber where it is drained into a sample jar, repeated pumping may be required to drain all the sample into the lower chamber. If preservation of the sample is necessary, this should be done after filtering.

2.7.6 Post Operation

After all samples are collected and preserved, the sampling equipment should be decontaminated prior to sampling another well. This will prevent cross-contamination of equipment and monitoring wells between locations.

- 1. Decontaminate all equipment.
- Replace sampling equipment in storage containers.
- 3. Prepare and transport water samples to the laboratory. Check sample documentation and make sure samples are properly packed for shipment.

2.7.7 Special Considerations for VOA Sampling

The proper collection of a sample for volatile organics requires minimal disturbance of the sample to limit volatilization and therefore a loss of volatiles from the sample.

Sample retrieval systems suitable for the valid collection of volatile organic samples are: positive displacement bladder pumps, gear driven submersible pumps, syringe samplers and bailers (Barcelona, 1984; Nielsen, 1985). Field conditions and other constraints will limit the choice of appropriate systems. The focus of concern must be to provide a valid sample for analysis, one which has been subjected to the least amount of turbulence possible.

The following procedures should be followed:

- Open the vial, set cap in a clean place, and collect the sample during the middle of the cycle. When collecting duplicates, collect both samples at the same time.
- 2. Fill the vial to just overflowing. Do not rinse the vial, nor excessively overfill it. There should be a convex meniscus on the top of the vial.
- 3. Check that the cap has not been contaminated (splashed) and carefully cap the vial. Place the cap directly over the top and screw down firmly. Do not overtighten and break the cap.

- 4 Invert the vial and tap gently. Observe vial for at least 10 seconds. If an air bubble appears, discard the sample and begin again. It is imperative that no entrapped air is in the sample vial.
- 5. Immediately place the vial in the protective foam sleeve and place into the cooler, oriented so that it is lying on its side, not straight up.
- 6. The holding time for VOAs is 7 days. Samples should be shipped or delivered to the laboratory daily so as not to exceed the holding time. Ensure that the samples remain at 4°C, but do not allow them to freeze.

2.8 CALCULATIONS

There are no calculations necessary to implement this procedure. However, if it is necessary to calculate the volume of the well, utilize the following equation:

Well volume =
$$nr^2h$$
 (cf) [Equation 1]

where:

n

radius of monitoring well (feet)
height of the water column (feet)
[This may be determined by subtracting the depth to water from the total depth of the well as measured from the same reference point.]

cf = conversion factor (gal/ft³) = 7.48

cf = conversion factor (gal/ft³) = 7.48 gal/ft³ [In this equation, 7.48 gal/ft³ is the necessary conversion factor.]

Monitoring wells are typically 2, 3, 4, or 6 inches in diameter. If you know the diameter of the monitoring well, there are a number of standard conversion factors which can be used to simplify the equation above.

The volume, in gallons per linear foot, for various standard monitoring well diameters can be calculated as follows:

$$v = nr^2$$
 (cf) [Equation 2)

where:

v = volume in gallons per linear foot

n = p

= radius of monitoring well (feet)

cf = conversion factor (7.48 gal/ft³)

For a 2-inch diameter well, the volume in gallons per linear foot can be calculated as follows:

```
v = nr<sup>2</sup> (cf) [Equation 2)
= 3.14 (l/12 ft)<sup>2</sup> 7.48 gal/ft<sup>3</sup>
= 0.1632 gal/ft
```

Remember that if you have a 2-inch diameter, well you must convert this to the radius in feet to be able to use the equation.

The volume in gallons per linear foot for the common size monitoring wells are as follows:

Well Diameter	v (volume in gal/ft.)
2 inches	0.1632
3 inches	0.3672
4 inches	0.6528
6 inches	1.4688

If you utilize the conversion factors above, Equation 1 should be modified as follows:

Well volume = (h)(v) [Equation 3)

where:

h = height of water column (feet)

v = volume in gallons per linear foot as calculated from Equation 2

2.9 QUALITY ASSURANCE/ QUALITY CONTROL

There are no specific quality assurance activities which apply to the implementation of these procedures. However, the following general QA procedures apply:

- All data must be documented on field data sheets or within site logbooks.
- All instrumentation must be operated in accordance with operating instructions as supplied by the manufacturer, unless

otherwise specified in the work plan. Equipment checkout and calibration activities must occur prior to sampling/operation and they must be documented.

2.10 DATA VALIDATION

This section is not applicable to this SOP.

2.11 HEALTH AND SAFETY

When working with potentially hazardous materials, follow U.S. EPA, OSHA and specific health and safety procedures. More specifically, depending upon the site-specific contaminants, various protective programs must be implemented prior to sampling the first well. The site health and safety plan should be reviewed with specific emphasis placed on the protection program planned for the well sampling tasks. Standard safe operating practices should be followed such as minimizing contact with potential contaminants in both the vapor phase and liquid matrix through the use of respirators and disposable clothing.

For volatile organic contaminants:

Avoid breathing constituents venting from the well.

- Pre-survey the well head-space with an FID/PID prior to sampling.
- ZZ If monitoring results indicate organic constituents, sampling activities may be conducted in Level C protection. At a minimum, skin protection will be afforded by disposable protective clothing.

Physical hazards associated with well sampling are:

- Lifting injuries associated with pump and bailer retrieval; moving equipment.
- Use of pocket knives for cutting discharge hose.
- Heat/cold stress as a result of exposure to extreme temperatures (may be heightened by protective clothing).
- Slip, trip, fall conditions as a result of pump discharge.
- Restricted mobility due to the wearing of protective clothing.

3.1 SCOPE AND APPLICATION

Soil gas monitoring provides a quick means of waste site evaluation. Using this method, underground contamination can be identified, and the source, extent, and movement of the pollutants can be traced.

This Standard Operating Procedure (SOP) outlines the methods used by EPA/ERT in installing soil gas wells; measuring organic levels in the soil gas using an HNU PI 101 Portable Photoionization Analyzer and/or other air monitoring devices; and sampling the soil gas using Tedlar bags, Tenax sorbent tubes, and SUMMA canisters.

3.2 METHOD SUMMARY

A 3/8-inch diameter hole is driven into the ground to a depth of 4 to 5 feet using a commercially available "slam bar". (Soil gas can also be sampled at other depths by the use of a longer bar or bar attachments.) A 1/Cinch O.D. stainless steel probe is inserted into the hole. The hole is then sealed at the top around the probe using modeling clay. The gas contained in the interstitial spaces of the soil is sampled by pulling the sample through the probe using an air sampling pump. The sample may be stored in Tedlar bags, drawn through sorbent cartridges, or analyzed directly using a direct reading instrument.

The air sampling pump is not used for SUMMA canister sampling of soil gas. Sampling is achieved by soil gas equilibration with the evacuated SUMMA canister. Other field air monitoring devices, such as the combustible gas indicator (MSA CGI/02 Meter, Model 260) and the organic vapor analyzer (Foxboro OVA, Model 128), can also be used depending on specific site conditions. Measurement of soil temperature using a temperature probe may also be desirable. Bagged samples are usually analyzed in a field laboratory using a portable Photovac GC.

Power driven sampling probes may be utilized when soil conditions make sampling by hand unfeasible (i.e., frozen ground, very dense clays, pavement, etc.). Commercially available soil gas sampling probes (hollow, 1/2inch O.D. steel probes) can be driven to the desired depth using a power hammer (e.g., Bosch Demolition Hammer). Samples can be drawn through the probe itself, or through Teflon tubing inserted through the probe and attached to the probe point. Samples are collected and analyzed as described above.

3.3 SAMPLE PRESERVATION, CONTAINERS, HANDLING, AND STORAGE

3.3.1 Tedlar Bag

Soil gas samples are generally contained in 1-L Tedlar bags. Bagged samples are best stored in coolers to protect the bags from any damage that may occur in the field or in transit. In addition, coolers ensure the integrity of the samples by keeping them at a cool temperature and out of direct sunlight. Samples should be analyzed as soon as possible, preferably within 24 to 48 hours.

3.3.2 Tenax Tube

Bagged samples can also be drawn into Tenax or other sorbent tubes to undergo lab GC/MS analysis. If Tenax tubes are to be utilized, special care must be taken to avoid contamination. Handling of the tubes should be kept to a minimum, and samplers must wear nylon or other lint-free gloves. After sampling, each tube should be stored in a clean, sealed culture tube; the ends packed with clean glass wool to protect the sorbent tube from breakage. The culture tubes should be kept cool and wrapped in aluminum foil to prevent any photodegradation of samples (see Section 3.7.4.).

3.3.3 SUMMA Canister

The SUMMA canisters used for soil gas sampling have a 6-L sample capacity and are certified clean by GC/MS analysis before being utilized in the field. After sampling is completed, they are stored and shipped in travel cases.

3.4 INTERFERENCES AND POTENTIAL PROBLEMS

3.4.1 HNU Measurements

A number of factors can affect the response of the HNU PI 101. High humidity can cause lamp fogging and decreased sensitivity. This can be significant when soil moisture levels are high, or when a soil gas well is actually in groundwater. High concentrations of methane can cause a downscale deflection of the meter. High and low temperature, electrical fields, FM radio transmission, and naturally occurring compounds, such as terpenes in wooded areas, will also affect instrument response.

Other field screening instruments can be affected by interferences. Consult the manufacturers' manuals.

3.4.2 Factors Affecting Organic Concentrations in Soil Gas

Concentrations in soil gas are affected by dissolution, adsorption, and partitioning. Partitioning refers to the ratio of component found in a saturated vapor above an aqueous solution to the amount in the solution; this can, in theory, be calculated using the Henry's Law constants. Contaminants can also be adsorbed onto inorganic soil components or "dissolved" in organic components. These factors can result in a lowering of the partitioning coefficient.

Soil "tightness" or amount of void space in the soil matrix, will affect the rate of recharging of gas into the soil gas well.

Existence of a high, or perched, water table, or of an impermeable underlying layer (such as a clay lens or layer of buried slag) may interfere with sampling of the soil gas. Knowledge of site geology is useful in such situations, and can prevent inaccurate sampling.

3.4.3 Soil Probe Clogging

A common problem with this sampling method is soil probe clogging. A clogged probe can be identified by using an in-line vacuum gauge or by listening for the sound of the pump laboring. This problem can usually be eliminated by using a wire cable to clear the probe (see procedure #3 in Section 3.7.1).

3.4.4 Underground Utilities

Prior to selecting sample locations, an underground utility search is recommended. The local utility companies can be contacted and requested to mark the locations of their underground lines. Sampling plans can then be drawn up accordingly. Each sample location should also be screened with a metal detector or magnetometer to verify that no underground pipes or drums exist.

3.5 EQUIPMENT/APPARATUS

3.5.1 Slam Bar Method

- slam bar (one per sampling team)
- soil gas probes, stainless steel tubing, 1/4-inch O.D., 5 foot length
- flexible wire or cable used for clearing the tubing during insertion into the well
- "quick connect" fittings to connect sampling probe tubing, monitoring instruments, and Gilian pumps to appropriate fittings on vacuum box
- modeling clay
- vacuum box for drawing a vacuum around Tedlar bag for sample collection (one per sampling team)
- Gilian pump Model HFS113A adjusted to approximately 3.0 L/min (one to two per sampling team)
- 1/4-inch Teflon tubing, 2 to 3 foot lengths, for replacement of contaminated sample line
- Tedlar bags, 1 liter, at least one bag per sample point
- soil gas sampling labels, field data sheets, logbook, etc.
- HNU Model PI 101, or other field air monitoring devices, (one per sampling team)
- ice chest, for carrying equipment and for protection of samples (two per sampling team)
- metal detector or magnetometer, for detecting underground utilities/pipes/drums (one per sampling team)
- Photovac GC, for field-lab analysis of bagged samples
- SUMMA canisters (plus their shipping cases) for sample, storage and transportation

3.5.2 Power Hammer Method

- Bosch demolition hammer
- 1/2inch O.D. steel probes, extensions, and points
- dedicated aluminum sampling points
- Teflon tubing, 1/Cinch O.D.
- "quick connect" fittings to connect sampling probe tubing, monitoring instruments, and Gilian pumps to appropriate fittings on vacuum box modeling clay
- vacuum box for drawing a vacuum around Tedlar bag for sample collection (one per sampling team)
- Gilian pump Model HFS113A adjusted to approximately 3.0 L/min (one to two per sampling team) 1/4-inch Teflon tubing, 2 to 3 foot lengths, for replacement of contaminated sample line
- Tedlar bags, 1 liter, at least one bag per sample point
- soil gas sampling labels, field data sheets, logbook, etc.
- HNU Model PI 101, or other field air monitoring devices, (one per sampling team)
- ice chest, for carrying equipment and for protection of samples (two per sampling team)
- metal detector or magnetometer, for detecting underground utilities/ pipes/drums (one per sampling team)
- Photovac GC, for field-lab analysis of bagged samples
- SUMMA canisters (plus their shipping cases) for sample, storage and transportation
- generator with extension cords
- high lift jack assembly for removing probes

3.6 REAGENTS

- HNU Systems Inc. Calibration Gas for HNU Model PI 101, and/or calibration gas for other field air monitoring devices
- deionized organic-free water, for decontamination
- methanol, HPLC grade, for decontamination
- ultra-zero grade compressed air, for field blanks

standard gas preparations for Photovac GC calibration and Tedlar bag spikes

3.7 PROCEDURES

3.7.1 Soil Gas Well Installation

- Initially, make a hole slightly deeper than the desired depth. For sampling up to 5 feet, use a 5-foot single piston slam bar. For deeper depths, use a piston slam bar with threaded 4foot-long extensions. Other techniques can be used, so long as holes are of narrow diameter and no contamination is introduced.
- 2. After the hole is made, carefully withdraw the slam bar to prevent collapse of the walls of the hole. Then insert the soil gas probe.
- 3. It is necessary to prevent plugging of the probe, especially for deeper holes. Place a metal wire or cable, slightly longer than the probe, into the probe prior to inserting into the hole. Insert the probe to full depth, then pull it up 3 to 6 inches, then clear it by moving the cable up and down. The cable is removed before sampling.
- 4. Seal the top of the sample hole at the surface against ambient air infiltration by using modeling clay molded around the probe at the surface of the hole.
- 5. If conditions preclude hand installation of the soil gas wells, the power driven system may be employed. Use the generator-powered demolition hammer to drive the probe to the desired depth (up to 12 feet may be attained with extensions). Pull the probe up 1 to 3 inches if the retractable point is used. No clay is needed to seal the hole. After sampling, retrieve the probe using the high lift jack assembly.
- 6. If semi-permanent soil gas wells are required, use the dedicated aluminum probe points. Insert these points into the bottom of the power-driven probe and attach it to the Teflon tubing. Insert the probe as in step 5. When the probe is removed, the point and Teflon tube remain in the hole, which may be scaled by backfilling with sand, bentonite, or soil.

3.7.2 Screening with Field Instruments

- 1. The **well** volume **must** be evacuated prior to sampling. Connect the Gilian pump, adjusted to 3.0 L/min, to the sample probe using a section of Teflon tubing as a connector. Turn the pump on, and a vacuum is pulled through the probe for approximately 15 seconds. A longer time is required for sample wells of greater depths.
- 2. After evacuation, connect the monitoring instrument(s) to the probe using a Teflon connector. When the reading is stable, or peaks, record the reading. For detailed procedures on HNU field protocol, see appendix B, and refer to the manufacturer's instructions.
- 3. Some readings may be above or below the range set on the field instruments. The range may be reset, or the response recorded as a figure greater than or less than the range. Consider the recharge rate of the well with soil gas when sampling at a different range setting.

3.7.3 Tedlar Bag Sampling

- 1. Follow step 1 in section 3.7.2 to evacuate well volume. If air monitoring instrument screening was performed prior to sampling, evacuation is not necessary.
- 2. Use the vacuum box and sampling train (Figure 3 in Appendix A) to take the sample. The sampling train is designed to minimize the introduction of contaminants and losses due to adsorption. All wetted parts are either Teflon or stainless steel. The vacuum is drawn indirectly to avoid contamination from sample pumps.
- 3. Place the Tedlar bag inside the vacuum box, and attach it to the sampling port. Attach the sample probe to the sampling port via Teflon tubing and a "quick connect" fitting.
- 4. Draw a vacuum around the outside of the bag, using a Gilian pump connected to the vacuum box evacuation port, via Tygon tubing and a "quick connect" fitting. The vacuum causes the bag to inflate, drawing the sample.

5. Break the vacuum by removing the Tygon line from the pump. Remove the bagged sample from the box and close valve. Label bag, record data on data sheets or in logbooks. Record the date, time, sample location ID, and the HNU, or other instrument reading(s) on sample bag label.

CAUTION: Labels should not be pasted directly onto the bags, nor should bags be labeled directly using a marker or pen. Inks and adhesive may diffuse through the bag material, contaminating the sample. Place labels on the edge of the bags, or tie the labels to the metal eyelets provided on the bags. Markers with inks containing volatile organics (i.e., permanent ink markers) should not be used.

3.7.4 Tenax Tube Sampling

Samples collected in Tedlar bags may be sorbed onto Tenax tubes for further analysis by GC/MS.

Additional Apparatus

- Syringe with a luer-lock tip capable of drawing a soil gas or air sample from a Tedlar bag onto a Tenax/CMS sorbent tube. The syringe capacity is dependent upon the volume of sample being drawn onto the sorbent tube.
- Adapters for fitting the sorbent tube between the Tedlar bag and the sampling syringe. The adapter attaching the Tedlar bag to the sorbent tube consists of a reducing union (I/Cinch to I/16-inch O.D. -- Swagelok cat. # SS-400-6-ILV or equivalent) with a length of I/4 inch O.D. Teflon tubing replacing the nut on the I/6-inch (Tedlar bag) side. A I/Cinch I.D. silicone O-ring replaces the ferrules in the nut on the I/Cinch (sorbent tube) side of the union.

The adapter attaching the sampling syringe to the sorbent tube consists of a reducing union (I/Cinch to I/16-inch O.D. --Swagelok Cat. # SS-4OO-6-ILV or equivalent) with a I/Cinch I.D. silicone O-ring replacing the ferrules in the nut on the I/4 inch (sorbent tube) side and the needle of a luer-lock syringe needle inserted into the I/16-inch side (held in place with a I/16-inch ferrule). The

luer-lock end of the needle can be attached to the sampling syringe. It is useful to have a luer-lock on/off valve situated between the syringe and the needle.

Two-stage glass sampling cartridge (1/4-inch O.D. x 1/&inch I.D. x 5 1/8 inch) contained in a flame-sealed tube (manufactured by Supelco Custom Tenax/Spherocarb Tubes or equivalent) containing two sorbent sections retained by glass wool:

Front section: 150 mg of Tenax-GC Back section: 150 mg of CMS (Carbonized Molecular Sieve)

Sorbent tubes may also be prepared in the lab and stored in either Teflon-capped culture tubes or stainless steel tube containers. Sorbent tubes stored in this manner should not be kept more than 2 weeks without reconditioning. (See SOP #2052 for Tenax/CMS sorbent tube preparation).

- Teflon-capped culture tubes or stainless steel tube containers for sorbent tube storage. These containers should be conditioned by baking at 120°C for at least 2 hours. The culture tubes should contain a glass wool plug to prevent sorbent tube breakage during transport. Reconditioning of the containers should occur between usage or after extended periods of disuse (i.e., 2 weeks or more).
- Nylon gloves or lint-free cloth. (Hewlett Packard Part # 8650-0030 or equivalent.)

Sample Collection

- 1. Handle sorbent tubes with care, using nylon gloves (or other lint-free material) to avoid contamination.
- 2. Immediately before sampling, break one end of the sealed tube and remove the Tenax cartridge. For in-house prepared tubes, remove cartridge from its container.
- 3. Connect the valve on the Tedlar bag to the sorbent tube adapter. Connect the sorbent tube to the sorbent tube adapter with the Tenax

(white granular) side of the tube facing the Tedlar bag.

- 4. Connect the sampling syringe assembly to the CMS (black) side of the sorbent tube. Fittings on the adapters should be very tight.
- **5.** Open the valve on the Tedlar bag.
- **6.** Open the on/off valve of the sampling syringe.
- 7. Draw a predetermined volume of sample onto the sorbent tube. (This may require closing the syringe valve, emptying the syringe and then repeating the procedure, depending upon the syringe capacity and volume of sample required.)
- 8. After sampling, remove the tube from the sampling train with gloves or a clean cloth. **Do not label or write on the Tenax/CMS tube.**
- 9. Place the sorbent tube in a conditioned stainless steel tube holder or culture tube. Culture tube caps should be sealed with Teflon tape.

Sample Labeling

Each sample tube container (not tube) must be labeled with the site name, sample station number, sample date, and sample volume.

Chain of custody forms must accompany all samples to the laboratory.

Quality Assurance

Before field use, a QA check should be performed on each batch of sorbent tubes by analyzing a tube with thermal desorption/cryogenic trapping GC/MS.

At least one blank sample must be submitted with each set of samples collected at a site. This trip blank must be treated the same as the sample tubes except no sample will be drawn through the tube.

Sample tubes should be stored out of UV light (i.e., sunlight) and kept on ice until analysis.

Samples should be taken in duplicate, when possible.

3.7.5 SUMMA Canister Sampling

- 1. Follow item 1 in step 3.7.2 to evacuate well volume. If HNU analysis was performed prior to taking a sample, evacuation is not necessary.
- 2. Attach a certified clean, evacuated 6-L SUMMA canister via the l/4-inch Teflon tubing.
- 3. Open the valve on SUMMA canister. The soil gas sample is drawn into the canister by pressure equilibration. The approximate sampling time for a 6-L canister is 20 minutes.
- 4. Site name, sample location, number, and date must be recorded on a chain of custody form and on a blank tag attached to the canister.

3.8 CALCULATIONS

3.8.1 Field Screening Instruments

Instrument readings are usually read directly from the meter. In some cases, the background level at the soil gas station may be subtracted:

Final Reading = Sample Reading - Background

3.8.2 Photovac GC Analysis

Calculations used to determine concentrations of individual components by Photovac GC analysis are beyond the scope of this SOP and are covered in ERT SOP #2109, Photovac GC Analysis for Soil, Water and Air/Soil Gas.

3.9 QUALITY ASSURANCE/ QUALITY CONTROL

3.9.1 Field Instrument Calibration

Consult the manufacturers' manuals for correct use and calibration of all instrumentation. The HNU should be calibrated at least once a day.

3.9.2 Gilian Model HFS113A Air Sampling Pump Calibration

Flow should be set at approximately 3.0 L/min;

accurate flow adjustment is not necessary. Pumps should be calibrated prior to bringing into the field.

3.9.3 Sample Probe Contamination

Sample probe contamination is checked between each sample by drawing ambient air through the probe via a Gilian pump and checking the response of the HNU PI 101. If HNU readings are higher than background, replacement or decontamination is necessary.

Sample probes may be decontaminated simply by drawing ambient air through the probe until the HNU reading is at background. More persistent contamination can be washed out using methanol and water, then air drying. Having more than one probe per sample team will reduce lag times between sample stations while probes are decontaminated.

3.9.4 Sample Train Contamination

The Teflon line forming the sample train from the probe to the Tedlar bag should be changed on a daily basis. If visible contamination (soil or water) is drawn into the sampling train, it should be changed immediately. When sampling in highly contaminated areas, the sampling train should be purged with ambient air, via a Gilian pump, for approximately 30 seconds between each sample. After purging, the sampling train can be checked using an HNU, or other field monitoring device, to establish the cleanliness of the Teflon line.

3.9.5 Field Blank

Each cooler containing samples should also contain one Tedlar bag of ultra-zero grade air, acting as a field blank. The field blank should accompany the samples in the field (while being collected) and when they are delivered for analysis. A fresh blank must be provided to be placed in the empty cooler pending additional sample collection. One new field blank per cooler of samples is required. A chain of custody form must accompany each cooler of samples and should include the blank that is dedicated to that group of samples.

3.9.6 Trip Standard

Each cooler containing samples should contain a Tedlar bag of standard gas to calibrate the analytical instruments (Photovac GC, etc.). This trip standard will be used to determine any changes in concentrations of the target compounds during the course of the sampling day (e.g., migration through the sample bag, degradation, or adsorption). A fresh trip standard must be provided and placed in each cooler pending additional sample collection. A chain of custody form should accompany each cooler of samples and should include the trip standard that is dedicated to that group of samples.

3.9.7 Tedlar Bag Check

Prior to use, one bag should be removed from each lot (case of 100) of Tedlar bags to be used for sampling and checked for possible contamination as follows: the test bag should be tilled with ultra-zero grade air; a sample should be drawn from the bag and analyzed via Photovac GC or whatever method is to be used for sample analysis. This procedure will ensure sample container cleanliness prior to the start of the sampling effort.

3.9.8 SUMMA Canister Check

From each lot of four cleaned SUMMA canisters, one is to be removed for a GC/MS certification check. If the canister passes certification, then it is re-evacuated and all four canisters from that lot are available for sampling.

If the chosen canister is contaminated, then the entire lot of four SUMMA canisters must be recleaned, and a single canister is re-analyzed by GC/MS for certification.

3.9.9 Options

Duplicate Samples

A minimum of 5% of all samples should be collected in duplicate (i.e., if a total of 100 samples arc to be collected, five samples should be duplicated). In choosing which samples to duplicate, the following criterion applies: if, after filling the first Tedlar bag, and, evacuating the well for 15 seconds, the second HNU (or other field monitoring device being used) reading matches or is close to (within 50%) the first reading, a duplicate sample may be taken.

Spikes

A Tedlar bag spike and Tenax tube spike may be desirable in situations where high concentrations of contaminants other than the target compounds are found to exist (landfills, etc.). The additional level of QA/QC attained by this practice can be useful in determining the effects of interferences caused by these non-target compounds. SUMMA canisters containing samples are not spiked.

3.10 DATA VALIDATION

For each target compound, the level of concentration found in the sample must be greater than three times the level (for that compound) found in the field blank which accompanied that sample to be considered valid. The same criteria apply to target compounds detected in the Tedlar bag pre-sampling contamination check.

3.11 HEALTH AND SAFETY

Because the sample is being drawn from underground, and no contamination is introduced into the breathing zone, soil gas sampling usually occurs in Level D, unless the sampling location is within the hot zone of a site, which requires Level B or Level C protection. However, to ensure that the proper level of protection is utilized, constantly monitor the ambient air using the HNU PI 101 to obtain background readings during the sampling procedure. As long as the levels in ambient air do not rise above background, no upgrade of the level of protection is needed.

Also, perform an underground utility search prior to sampling (see section 3.4.4). When working with potentially hazardous materials, follow U.S. EPA, OSHA, and specific health and safety procedures.

4.0 MONITORING WELL INSTALLATION: SOP #2150

4.1 SCOPE AND APPLICATION

The purpose of this Standard Operating Procedure (SOP) is to provide an overview of the methods used for monitoring well installation. Monitoring well installation creates a permanent access for the collection of samples to determine groundwater quality and the hydrogeologic properties of the aquifer in which the contaminants exist. Such wells should not alter the medium which is being monitored.

The most commonly used drilling methods are: (1) hollow-stem augers, (2) cable tool drills, and (3) rotary drills. Rotary drilling can be divided into a mud rotary or air rotary method.

4.2 METHOD SUMMARY

There is no ideal monitoring well installation method for all conditions; therefore, hydrogeologic conditions at the site and project objectives must be considered before deciding which drilling method to use.

4.2.1 Hollow-Stem Augering

Hollow-stem augering is fast and relatively less expensive than cable tool or rotary drilling methods. It is possible to drill several hundred feet of borehole per day in unconsolidated formations.

4.2.2 Cable Tool Drilling

Cable tool drilling method involves lifting and dropping a heavy, solid chisel-shaped bit, suspended on a steel cable. This bit pounds a hole through soil and rock. Temporary steel casing is used while drilling to keep the hole open and to isolate strata. The temporary casing is equipped with a drive shoe, which is attached to the lower end, and which aids the advancement of the casing by drilling out a slightly larger diameter borehole than the hole made by the drill bit alone.

Water is sometimes used when drilling above the saturated zone to reduce dust and to form a slurry with the loosened material. This facilitates removal of cuttings using a bailer or a sand pump. Potable

water or distilled/deionized water should be used to prevent the introduction of contamination into the borehole.

4.2.3 Rotary Drilling

Mud Rotary Method

In the mud rotary method, the borehole is advanced by rapid rotation of the drill bit, which cuts and breaks the material at the bottom of the hole into smaller pieces. Cuttings are removed by pumping drilling fluid (water, or water mixed with bentonite) down through the drill rods and bit, and up the armulus between the borehole and the drill rods. The drilling fluid also serves to cool the drill bit and prevent—the borehole from collapsing in unconsolidated formations.

Air Rotary Method

The air rotary method is the same as the mud rotary except that compressed air is pumped down the drill rods and returns with the drill cuttings up through the annulus. Air rotary method is generally limited to consolidated and semi-consolidated formations. Casing is sometimes used to prevent cavings in semi-consolidated formations. The air must be filtered to prevent introduction of contamination into the borehole.

4.3 SAMPLE PRESERVATION, CONTAINERS, HANDLING, AND STORAGE

Often, a primary object of the drilling program is to obtain representative lithologic or environmental samples. Lithologic samples are taken in order to determine the geologic or hydrogeologic regime at a site. The most common techniques for retrieving lithologic samples in unconsolidated formations are described below.

Split spoon sampling, carried out continuously or at discrete intervals during drilling, is used to make a field description of the sample and create a log of each boring.

- Shelby tube sampling, is used when an undisturbed sample is required from clayey or silty soils, especially for geotechnical evaluation or chemical analysis.
- Cuttings description is used when a general lithologic description and approximate depths are sufficient.

The most common techniques for retrieving lithologic sampling in consolidated formations are described below.

- Rock coring is carried out continuously or at discrete intervals during drilling and enables the geologist to write a field description of the sample, create a log of each boring, and map occurrences and orientation of fractures.
- Cuttings description is used when a general lithologic description and approximate depths are sufficient.

4.4 INTERFERENCES AND POTENTIAL PROBLEMS

Table 3 on page 27 displays the advantages and disadvantages of the various drilling techniques.

4.5 EQUIPMENT/APPARATUS

The drilling contractor will provide all operational equipment for the drilling program which is outlined. The geologist should bring:

- well log sheets
- metal case (container for well logs)
- ruler
- zz depth sounder
- water level indicator
- all required health and safety gear
- sample collection jars
- zz trowels
- description aids (Munsell, grain size charts, etc.)

4.6 REAGENTS

No chemical reagents are used in this procedure. Decontamination of drilling equipment should follow ERT SOP #2006, Sampling Equipment Decontamination and the site-specific work plan.

4.7 PROCEDURES

4.7.1 Preparation

The planning, selection and implementation of any monitoring well installation program should include the following steps.

- Review existing data on site geology and hydrogeology including publications, air photos, water quality data, and existing maps. These may be obtained from local, state, or federal agencies.
- 2. Visit the site to observe field geology and potential access problems for drill rig, to establish water supply, and drill equipment and materials storage area.
- 3. Prepare site safety plan.
- 4. Define project objectives; select drilling, well development, and sampling methods.
- Select well construction materials including well construction specifications (i.e., casing and screen materials, casing and screen diameter, screen length and screen interval, filter pack and screen size).
- 6. Determine need for containing drill cuttings/fluids and their disposal.
- 7. Prepare work plan including all of the above.
- 8. Prepare and execute the drilling contract.
- 9. Implement the drilling program.
- Prepare the final report, including background data, project objective, field procedure, well construction data including well logs and well construction.

All drilling and well installation programs must be planned and supervised by a professional geologist/hydrogeologist.

4.7.2 Field Preparation

1. Prior to the mobilization of the drill rig,

Table 3: Advantages and Disadvantages of Various Drilling Techniques

Drilling Type	Advantages	Disadvantages
Auger	Allows sampling from different strata during drilling. Less potential for cross-contamination between strata than in other techniques. Large diameter borehole may be drilled for multiple-well completion. Less well development is generally needed than in mud rotary because of the relatively large diameter borehole, the ability to emplace a large and effective gravel pack, and because no drilling fluids are introduced into the borehole.	 Very slow or impossible in coarse materials such as cobbles and boulders. Cannot drill hard rock formations and is generally not suited for wells deeper than 100 feet. Not good in caving formations. Potential for disturbing large volume of subsurface materials around the borehole; therefore affecting local permeabilities and creating annular channels for contaminant movement between different strata.
Cable Tool	Allows for easy and accurate detection of the water table. Driven casing seals off formation, minimizing the threat of cross-contamination in pollution investigation. Especially successful for drilling in glacial till.	Extremely slow rate of drilling. Can lose casing in deep wells.
Mud Rotary	Quite fast, more than 100 feet of borehole advancement per day is possible. Geophysical logs such as resistivity (which must be run in an uncased borehole) can be run before well construction.	Potential cross-contamination of strata exposed to the circulating drilling fluid during drilling. Difficulty in removing mud residues during well development. Drilling mud may alter the groundwater chemistry by binding metals, sorbing organic compounds and by altering pH, cation exchange capacity and chemical oxidation demand of native fluids. Drilling mud may change local permeability of the formation.
Air Rotary	Like mud rotary method, more than 100 feet of borehole advancement a day is possible. Sampling different strata during drilling is possible if temporary casing is advanced.	In contaminated formations, the use of high pressure air may pose a significant hazard to the drill crew due to rapid transport of contaminated material up the borehole during drilling. Introduction of air to ground water could reduce concentration of volatile organic compounds locally.

thoroughly decontaminate the rig and all associated equipment to remove all oil, grease, mud, etc.

- 2. Before drilling each boring, steam-clean and rinse all the "down-the-hole" drill equipment with potable water to minimize cross-contamination. Special attention should be given to the thread section of the casings, and to the drill rods. All drilling equipment should be steam-cleaned at completion of the project to ensure that no contamination is transported to or from the sampling site.
- 3. Record lithologic descriptions and all field measurements and comments on the well log form (Appendix C). Include well construction diagrams on the well log form for each well installed. At a minimum, the well construction information should show depth from surface grade, the bottom of the boring, the screened interval, casing material, casing diameter, gravel pack location, grout seal and height of riser pipe above the ground. Also record the actual compositions of the grout and seal on the well log form.

4.7.3 Well Construction

The most commonly used casing materials include stainless steel, polyvinyl chloride (PVC) and Teflon. Monitoring wells are constructed with casings and materials that are resistant to the subsurface environment. The selection of well construction material is based on the material's long-term interaction with the contaminated groundwater. Construction materials should not cause an analytical bias in the interpretation of the chemical analysis of the water samples.

Well casing material should also be judged from a structural standpoint. Material should be rigid and nonporous, with a low surface-area-to-water ratio in the wellbore relative to the formation materials (U.S. EPA, 1987).

1. Fill the annular space between the well screen and the boring with a uniform gravel/sand pack to serve as a filter media. For wells deeper than approximately 50 feet, or when recommended by the site geologist, emplace the sand pack using a tremie pipe (normally consisting of a 1.25inch PVC or steel pipe). Pump sand slurry composed of sand and

potable water through the tremie pipe into the annulus throughout the entire screened interval, and over the top of the screen. It is necessary to pump sufficient sand/gravel slurry to cover the screen after the sand/gravel pack has settled and become dense.

- 2. Determine the depth of the top of the sand using the tremie pipe, thus verifying the thickness of the sand pack. Add more sand to bring the top of the sand pack to approximately 2-3 feet above the top of the well screen. Under no circumstances should the sand pack extend into any aquifer other than the one to be monitored. In most cases, the well design can be modified to allow for a sufficient sand pack without threat of crossflow between producing zones through the sand pack.
- In materials that will not maintain an open hole, withdraw the temporary or outer casing gradually during placement of sand pack/grout to the extent practical.

For example, after filling 2 feet with sand pack, the outer casing should be withdrawn 2 feet. This step of placing more gravel and withdrawing the outer casing should be repeated until the level of the sand pack is approximately 3 feet above the top of the well screen. This ensures that there is no locking of the permanent (inner) casing in the outer casing.

 Emplace a bentonite seal, composed of pellets, between the sand pack and grout to prevent infiltration of cement into the filter pack and the well screen.

These pellets should have a minimum purity of 90% montmorillonite clay, and a minimum dry bulk density of 75 lb/ft³ for ½ inch pellets, as provided by American Colloid, or equivalent. Bentonite pellets shall be poured directly down the annulus.

Care must be taken to avoid introducing pellets into the well bore. A cap placed over the top of the well casing before pouring the bentonite pellets from the bucket will prevent this. To ensure even application, pour the pellets from different points around the casing. To avoid bridging of pellets, they should not be introduced at a rate faster than they can settle. A tremie pipe may be used to redistribute and

level out the top of the seal,

- 5. If using a slurry of bentonite as an annular seal, prepare it by mixing powdered or granular bentonite with potable water. The slurry must be of sufficiently high specific gravity and viscosity to prevent its displacement by the grout to be emplaced above it. As a precaution, regardless of depth, and depending on fluid viscosity, add a few handfuls of bentonite pellets to solidify the bentonite slurry surface.
- 6. Place a mixture of cement and bentonite grout from the top of the bentonite seal to the ground surface.

Only Type I or II cement without accelerator additives may be used. An approved source of potable water must be used for mixing grouting materials. The following mixes are acceptable:

- Neat cement, a maximum of 6 gallons of water per 94-pound bag of cement
- Granular bentonite, 1.5 pounds of bentonite per 1 gallon of water
- Cement-bentonite, 5 pounds of pure bentonite per 94-pound bag of cement with 7-8 gallons of water; 13-14 pounds weight, if dry mixed
- Cement-bentonite, 6 to 8 pounds of pure bentonite per 94-pound bag of cement with 8-10 gallons of water, if water mixed
- Non-expandable cement, mixed at 7.5 gallons of water to 1/2 teaspoon of aluminum hydroxide, 94 pounds of neat cement (Type I) and 4 pounds of bentonite
- Non-expandable cement, mixed at 7 gallons of water to 1/2 teaspoon of aluminum hydroxide, 94 pounds of neat cement (Type I and Type II)
- 7. Pump grout through a tremie pipe to the bottom of the open annulus until undiluted grout flows from the annulus at the ground surface.
- 8. In materials that will not maintain an open hole, the temporary steel casing should be withdrawn in a manner that prevents the level

- of grout from dropping below the bottom of the casing.
- Additional grout may be added to compensate for the removal of the temporary casing and the tremie pipe to ensure that the top of the grout is at or above ground surface.
- 10. Place the protective casing. Protective casings should be installed around all monitoring wells. Exceptions are on a case-by-case basis. The minimum elements in the protection design include:
 - A protective steel cap to keep precipitation out of the protective casing, secured to the casing by padlocks.
 - A 5-foot-minimum length of black iron or galvanized pipe, extending about 1.5 to 3 feet above the ground surface, and set in cement grout. The pipe diameter should be 8 inches for 4-inch wells, and 6 inches for 2-inch wells (depending on approved borehole size). A 0.5-inch drain hole near ground level is permitted.
 - The installation of guard posts in addition to the protective casing, in areas where vehicular traffic may pose a hazard. These guard posts consist of 3-inch diameter steel posts or tee-bar driven steel posts. Groups of three are radially located 4 feet around each well 2 feet below and 4 feet above ground surface, with flagging in areas of high vegetation. Each post is cemented inplace.
 - A flush mount of protective casing may also be used in areas of high traffic or where access to other areas would be limited by a well with stickup.

After the grout sets (about 48 hours), fill any depression due to settlement with a grout mix similar to that described above.

4.0 CALCULATIONS

To maintain an open borehole using sand or water rotary drilling, the drilling fluid must exert a pressure greater than the formation pore pressure. Typical pore pressure for an unconfined aquifer is 0.433 psi/ft and for a confined aquifer is 0.465 psi/ft.

The calculation for determining the hydrostatic pressure of the drilling fluid is:

Hydrostatic Pressure (psi) = Fluid Density (lb/gal) x Height of Fluid Column (ft) x 0.052

The minimum grout volume necessary to grout a well can be calculated using:

Grout Vol (ft³) = Vol of Borehole (f³) -
Vol of Casing (ft³) = L (
$$r_b^2 - r_c^2$$
)

where:

r_B = radius of boring (ft)
 rc = radius of casing (ft)
 L = length of borehole to be grouted (ft)

4.9 QUALITY ASSURANCE/ QUALITY CONTROL

There are no specific quality assurance activities which apply to the implementation of these procedures.

However, the following general QA procedures apply:

- All data must be documented on standard well completion forms, field data sheets or within field/site logbooks.
- All instrumentation must be operated in accordance with operating instructions as supplied by the manufacturer, unless otherwise specified in the work plan. Equipment checkout and calibration activities must occur prior to sampling/operation and they must be documented.

4.10 DATA VALIDATION

This section is not applicable to this SOP.

4.11 HEALTH AND SAFETY

When working with potentially hazardous materials, follow U.S. EPA, OSHA, and specific health and safety procedures.

5.0 WATER LEVEL MEASUREMENT: SOP #2151

5.1 **SCOPE AND APPLICATION**

The purpose of this Standard Operating Procedure (SOP) is to set guidelines for the determination of the depth to water in an open borehole, cased borehole, monitoring well or piezometer.

Generally, water level measurements from boreholes, piezometers, or monitoring wells are used to construct water table or potentiometric surface maps. Therefore, all water level measurements at a given site should be collected within a 24-hour period. Certain situations may necessitate that all water level measurements be taken within a shorter time interval. These situations may include:

- the magnitude of the observed changes between wells appears too large
- atmospheric pressure changes
- aquifers which are tidally influenced
- aquifers affected by river stage, impoundments, and/or unlined ditches
- aquifers stressed by intermittent pumping of production wells
- aquifers being actively recharged due to precipitation events

5.2 METHOD SUMMARY

A survey mark should be placed on the casing for use as a reference point for measurement. Many times the lip of the riser pipe is not flat. Another measuring reference should be located on the grout apron. The measuring point should be documented in the site logbook and on the groundwater level data form (see Appendix C).

Water levels in piezometers and monitoring wells should be allowed to stabilize for a minimum of 24 hours after well construction and development, prior to measurement. In low yield situations, recovery may take longer.

Working with decontaminated equipment, proceed from the least to the most contaminated wells. Open the well and monitor headspace with the appropriate monitoring instrument to determine the presence of volatile organic compounds. Lower the water level measurement device into the well until water surface or bottom of casing is encountered. Measure distance from water surface to the reference point on the well casing and record in the site logbook and/or groundwater level data form. Remove all downhole equipment, decontaminate as necessary, and replace well casing cap.

5.3 SAMPLE PRESERVATION, CONTAINERS, HANDLING AND STORAGE

This section is not applicable to this SOP.

5.4 INTERFERENCES AND POTENTIAL PROBLEMS

- The chalk used on steel tape may contaminate the well.
- Cascading water may obscure the water mark or cause it to be inaccurate.
- Many types of electric sounders use metal indicators at 5-foot intervals around a conducting wire. These intervals should be checked with a surveyor's tape to ensure accuracy.
- If there is oil present on the water, it can insulate the contacts of the probe on an electric sounder or give false readings due to thickness of the oil. Determining the thickness and density of the oil layer may be warranted, in order to determine the correct water level.
- Turbulence in the well and/or cascading water can make water level determination difficult with either an electric sounder or steel tape.

An airline measures drawdown during pumping. It is only accurate to 0.5 foot unless it is calibrated for various "drawdowns".

5.5 EQUIPMENT/APPARATUS

There are a number of devices which can be used to measure water levels, such as steel tape or airlines. The device should be adequate to attain an accuracy of 0.01 feet.

The following equipment is needed to measure water levels:

- air monitoring equipment
- water level measurement device
- electronic water level indicator
- metal tape measure
- airline
- steel tape
- ∠ chalk
- ∠ ruler
- notebook
- ≠ paper towels
- decontamination solution and equipment
- groundwater level data forms

5.6 REAGENTS

No chemical reagents are used in this procedure, with the exception of decontamination solutions. Where decontamination of equipment is required, refer to ERT SOP #2006, Sampling Equipment Decontamination and the site-specific work plan.

5.7 PROCEDURES

57.1 Preparation

- 1. Determine the extent of the sampling effort, the sampling methods to be employed, and which equipment and supplies are needed.
- 2. Obtain necessary sampling and monitoring equipment.
- 3. Decontaminate or preclean equipment, and ensure that it is in working order.
- 4. Prepare scheduling and coordinate with staff,

- clients, and regulatory agency, if appropriate.
- Perform a general site survey prior to site entry in accordance with the site-specific health and safety plan.
- 6. Identify and mark all sampling locations.

5.7.2 Procedures

- 1. Make sure water level measuring equipment is in good operating condition.
- 2. If possible and where applicable, start at those wells that are least contaminated and proceed to those wells that are most contaminated.
- 3. Clean all equipment entering the well by the following decontamination procedure:
 - Triple rinse equipment with deionized water.
 - Wash equipment with an Alconox solution followed by a deionized water rinse.
 - Rinse with an approved solvent (e.g., methanol, isopropyl alcohol, acetone) as per the work plan, if organic contamination is suspected.
 - Place equipment on clean surface such as a Teflon or polyethylene sheet.
- 4. Remove locking well cap, note location, time of day, and date in site notebook or an appropriate groundwater level data form.
- 5. Remove well casing cap.
- If required by site-specific condition, monitor headspace of well with PID or PID to determine presence of volatile organic compounds and record in site logbook.
- 7. Lower electric water level measuring device or equivalent (i.e., permanently installed tranducers or airline) into the well until water surface is encountered.
- 8. Measure the distance from the water surface to the reference measuring point on the well casing or protective barrier post and record in the field logbook. In addition, note that the

water level measurement was from the top of the steel casing, top of the PVC riser pipe, from the ground surface, or from some other position on the well head.

- 9. The groundwater level data form in Appendix C should be completed as follows:
 - site name
 - ø logger name: person taking field notes
 - date: the date when the water levels are being measured
 - location: monitor well number and physical location
 - time: the military time at which the water level measurement was recorded
 - depth to water: the water level measurement in feet, or in tenths or hundreds of feet, depending on the equipment used
 - comments: any information the field personnel feels to be applicable
 - measuring point: marked measuring point on PVC riser pipe, protective steel casing or concrete pad surrounding well casing from which all water level measurements for individual wells should be measured. This provides consistency in future water level measurements.
- 10. Measure total depth of well (at least twice to confirm measurement) and record in site notebook or on log form.
- 11. Remove all downhole equipment, replace well casing cap and lock steel caps.
- 12. Rinse all downhole equipment and store for transport to next well.
- 13. Note any physical changes such as erosion or cracks in protective concrete pad or variation in total depth of well in field notebook and on field data sheets.
- 14. Decontaminate all equipment as outlined in Step 3 above.

5.8 CALCULATIONS

To determine groundwater elevation above mean sea level, use the following equation:

Ew = E-D

where:

Ew = Elevation of water above mean sea

E = Elevation above sea level at point of measurement

D = Depth to water

5.9 QUALITY ASSURANCE/ QUALITY CONTROL

The following general quality assurance procedures apply:

- All data must be documented on standard chain of custody forms, field data sheets or within personal/site logbooks.
- All instrumentation must be operated in accordance with operating instructions as supplied by the manufacturer, unless otherwise specified in the work plan. Equipment checkout and calibration activities must occur prior to sampling/operation, and they must be documented.
- Each well should be tested at least twice in order to compare results.

5.10 DATA VALIDATION

This section is not applicable to this SOP.

5.11 HEALTH AND SAFETY

When working with potentially hazardous materials, follow U.S. EPA, OSHA, and specific health and safety procedures.

6.0 WELL DEVELOPMENT: SOP #2156

6.1 SCOPE AND APPLICATION

The purpose of monitoring well development is to ensure removal of fines from the vicinity of the well screen. This allows free flow of water from the formation into the well and also reduces the turbidity of the water during sampling events. The most common well development methods are: surging, jetting, and overpumping.

Surging involves raising and lowering a surge block or surge plunger inside the well. The resulting motion surges water into the formation and loosens sediment to be pulled from the formation into the well. Occasionally, sediment must be removed from the well with a sand bailer to prevent sand locking of the surge block. This method may cause the sand pack around the screen to be displaced to a degree that damages its value as a filtering medium. For example, channels or voids may form near the screen if the filter pack sloughs away during surging (Keely and Boateng, 1987).

Jetting involves lowering a small diameter pipe into the well to a few feet above the well screen, and injecting water or air through the pipe under pressure so that sediments at the bottom are geysered out the top of the well. It is important not to jet air or water directly across the screen. This may cause fines in the well to be driven into the entrance of the screen openings thereby causing blockages.

Overpumping involves pumping at a rate rapid enough to draw the water level in the well as low as possible, and allowing it to recharge. This process is repeated until sediment-free water is produced. Overpumping is not as vigorous as surging and jetting and is probably the most desirable for monitoring well development.

6.2 METHOD SUMMARY

Development of a well should occur as soon as practical after installation, but not sooner than 48 hours after grouting is completed, if a rigorous well development is being used. If a less rigorous method, such as bailing, is used for development, it may be initiated shortly after installation. The main

concern is that the method being used for development does not interfere with allowing the grout to set.

Open the monitoring well, take initial measurements (e.g. head space air monitoring readings, water level, well depth, pH, temperature, and specific conductivity) and record results in the site logbook. Develop the well by the appropriate method (i.e., overpumping jetting, or surging) to accommodate site conditions and project requirements. Continue until the developed water is clear and free of sediment. Containerize all discharge water from known or suspected contaminated areas. Record final measurements in the logbook. Decontaminate equipment as appropriate prior to use in the next well.

6.3 SAMPLE PRESERVATION, CONTAINERS, HANDLING, AND STORAGE

This section is not applicable to this Standard Operating Procedure (SOP).

6.4 INTERFERENCES AND POTENTIAL PROBLEMS

The following interferences or problems may occur during well development:

- The possibility of disturbing the filter pack increases with surging and jetting well development methods.
- The introduction of external water or air by jetting may alter the hydrochemistry of the aquifer.

6.5 EQUIPMENT/APPARATUS

The type of equipment used for well development is dependent on the diameter of the well. For example, submersible pumps cannot be used for well development unless the wells are 4 inches or greater in diameter, because the smallest submersible pump has a 3 1/4 inch O.D.

In general, the well should be developed shortly after it is drilled. Most drilling rigs have air compressors or pumps that may be used for the development process.

6.6 REAGENTS

No chemical reagents are used in this procedure except for decontamination solutions. For guidelines on equipment decontamination, refer to ERT SOP #2006, Sampling Equipment Decontamination and the site-specific work plan.

6.7 PROCEDURES

6.7.1 Preparation

- 1. Coordinate site access and obtain keys to the monitoring well security cap locks.
- 2. Obtain information on each well to be developed (i.e., drilling, method, well diameter, depth, screened interval, anticipated contaminants, etc.).
- 3. Obtain a water level meter, air monitoring equipment, materials for decontamination, pH and electrical conductivity meters, a thermometer, and a stopwatch.
- 4. Assemble containers for temporary storage of water produced during well development. Containers must be structurally sound, compatible with anticipated contaminants, and easy to manage in the field. The use of truck-mounted tanks may be necessary in some cases; alternately, a portable water treatment unit (e.g. activated carbon) may be used to decontaminate the purge water.

6.7.2 Operation

The development should be performed as soon as practical after the well is installed, but no sooner than 48 hours after grouting is completed. Dispersing agents, acids, or disinfectants should not be used to enhance development of the well.

1. Assemble necessary equipment on a plastic sheet around the well.

- Record pertinent information in field logbook (personnel, time, location ID, etc.).
- 3. Open monitoring well, and take air monitoring readings at the top of casing and in the breathing zone as appropriate.
- 4. Measure depth to water and the total depth of the monitoring well from the same datum point.
- 5. Measure the initial pH, temperature, and specific conductivity of the water and record in the logbook.
- 6. Develop the well until the water is clear and appears to be free of sediment. Note the initial color, clarity and odor of the water.
- 7. All water produced by development in contaminated or suspected contaminated areas must be containerized or treated. Clearly label each container with the location ID. Determination of the appropriate disposal method will be based on the fast round of analytical results from each well.
- 8. No water should be added to the well to assist development without prior approval by the site geologist. If a well cannot be cleaned of mud to produce formation water because the aquifer yields insufficient water, small amounts of potable water may be injected to clean up this poorly yielding well. This may be done by dumping in buckets of water. When most of the mud is out, continue development with formation water only. It is essential that at least live times the amount of water injected must be produced back from the well in order to ensure that all injected water is removed from the formation.
- 9. Note the final color, clarity and odor of the water.
- 10. Measure the final pH, temperature and specific conductance of the water and record in the field logbook.
- 11. Record the following data in the field logbook:
 - well designation (location ID)
 - ∠ date(s) of well installation
 - ∠ date(s) and time of well development
 - static water level before and after

development

- ex quantity of water removed and time of removal
- type and size/capacity of pump and/or bailer used
- description of well development techniques used

6.7.3 Post Operation

- 1. Decontaminate all equipment.
- 2. Store containers of purge water produced during development in a safe and secure area.
- 3. After the first round of analytical results have been received, determine and implement the appropriate purge water disposal method.

6.8 CALCULATIONS

There are no calculations necessary to implement this procedure. However, if it is necessary to calculate the volume of the well, utilize the following equation:

Well volume =
$$nr^2h(cf)$$
 [Equation 1]

where:

n = pi

r = radius of monitoring well (feet)

h = height of the water column (feet)
[This may be determined by
subtracting the depth to water
from the total depth of the well as
measured from the same reference
point.]

cf = conversion factor (gal/ft³) = 7.48 gal/ft³ [In this equation, 7.48 gal/ft³ is the necessary conversion factor, because 7.48 gallons of

water occupy 1 ft³]

Monitoring wells are typically 2 inches, 3 inches, 4 inches, or 6 inches in diameter. If the diameter of the monitoring well is known, a number of standard conversion factors can be used to simplify the equation above.

The volume, in gallons per linear foot, for various standard monitoring well diameters can be calculated as follows:

 $v = nr^2(cf)$ [Equation 2]

where:

v = volume in gallons per linear foot

 $n = p_i$

r = radius of monitoring well (feet) cf = conversion factor (7.48 gal/ft³)

For a 2-inch diameter well, the volume per linear foot can be calculated as follows:

 $V = nr^{2}(cf)$ [Equation 21 = 3.14 (l/12 ft)² 7.48 gal/f?

= 0.1632 gal/ft

Remember that if you have a 2-inch diameter well, you must convert this to the radius in feet to be able to use the equation.

The volume per linear foot for monitoring wells of common size are as follows:

Well diameter	v (volume in gal/ft.)
2-inch	0.1632
3-inch	0.3672
4-inch	0.6528
6-inch	1.4688

If you utilize the factors above, Equation 1 should be modified as follows:

Well volume = h(v) [Equation 3]

where:

h = height of water column (feet) v = volume in gallons per linear foot from Equation 2

6.9 QUALITY ASSURANCE/ QUALITY CONTROL

There are no specific quality assurance activities which apply to the implementation of these procedures. However, the following general QA procedures apply:

- All data must be documented on standard chain of custody forms, field data sheets or personal/site logbooks.
- All instrumentation must be operated in accordance with operating instructions as

supplied by the manufacturer, unless otherwise specified in the work plan. Equipment checkout and calibration activities must occur prior to sampling/operation and they must be documented.

6.10 DATA VALIDATION

This section is not applicable to this SOP

6.11 HEALTH AND SAFETY

7.0 CONTROLLED PUMPING TEST: SOP #2157

7.1 SCOPE AND APPLICATION

The most reliable and commonly used method of determining aquifer characteristics is by controlled aquifer pumping tests. Groundwater flow varies in space and time and depends on the hydraulic properties of the rocks and the boundary conditions imposed on the groundwater system. Pumping tests provide results that are more representative of aquifer characteristics than those predicted by slug or bailer tests. Pumping tests require a greater degree of activity and expense, however, and are not always justified for all levels of investigation. For example, slug tests may be acceptable at the reconnaissance level whereas pumping tests are usually performed as part of a feasibility study in support of designs for aquifer remediation.

Aquifer characteristics which may be learned using pumping tests include hydraulic conductivity (K), transmissivity (T), specific yield (Sy) for unconfined aquifers, and storage coefficient (S) for confined aquifers. These parameters can be determined by graphical solutions and computerized programs. This Standard Operating Procedure (SOP) outlines the protocol for conducting controlled pumping tests.

7.2 METHOD SUMMARY

It is desirable to monitor pre-test water levels at the test site for about 1 week prior to performance of the pump test. This information allows for the determination of the barometric efficiency of the aquifer, as well as noting changes in head, due to recharging or pumping in the area adjacent to the well. Prior to initiating the long term pump test, a step test is conducted to estimate the greatest flow rate that may be sustained by the pump well.

After the pumping well has recovered from the step test, the long term pumping test begins. At the beginning of the test, the discharge rate is set as quickly and accurately as possible. The water levels in the pumping well and observation wells are recorded accordingly with a set schedule. Data is entered on the Pump/Recovery Test Data Sheet (Appendix C). The duration of the test is determinated by project needs and aquifer

properties, but rarely goes beyond 3 days or until water levels become constant.

7.3 SAMPLE PRESERVATION, CONTAINERS, HANDLING, AND STORAGE

This section is not applicable to this SOP.

7.4 INTERFERENCES AND POTENTIAL PROBLEMS

Interferences and potential problems include:

- atmospheric conditions
- impact of local potable wells
- compression of the aquifer due to trains, heavy traffic, etc.

7.5 EQUIPMENT/APPARATUS

- tape measure (subdivided into tenths of feet)
- submersible pump
- water pressure transducer
- electric water level indicator
- weighted tapes
- steel tape (subdivided into tenths of feet)
- generator
- electronic data-logger (if transducer method is used)
- watch or stopwatch with second hand
- semilogarithmic graph paper (if required)
- water proof ink pen and logbook
- thermometer
- appropriate references and calculator
- a barometer or recording barograph (for tests conducted in confined aquifers)
- heat shrinks
- electrical tape
- flashlights and lanterns
- pH meter
- conductivity meter
- discharge pipe
- flow meter

7.6 REAGENTS

No chemical reagents are used for this procedure; however, decontamination solutions may be necessary. If decontamination of equipment is required, refer to ERT SOP #2006, Sampling Equipment Decontamination and the site-specific work plan.

7.7 PROCEDURES

7.7.1 Preparation

- 1. Determine the extent of the sampling effort, the sampling methods to be employed, and which equipment and supplies are needed.
- 2. Obtain necessary sampling and monitoring equipment.
- 3. Decontaminate or preclean equipment, and ensure that it is in working order.
- 4. Prepare scheduling and coordinate with staff, clients, and regulatory agency, if appropriate.
- Perform a general site survey prior to site entry in accordance with the site-specific health and safety plan.
- 6. Identify and mark all sampling locations.

7.7.2 Field Preparation

- 1. Review the site work plan and become familiar with information on the wells to be tested.
- Check and ensure the proper operation of all field equipment. Ensure that the electronic data-logger is fully charged, if appropriate. Test the electronic data-logger using a container of water. Always bring additional transducers in case of malfunctions.
- 3. Assemble a sufficient number of field data forms to complete the field assignment.
- 4. Develop the pumping well prior to testing, per ERT SOP #2156, Well Development.
- Provide an orifice, weir, flow meter, container or other type of water measuring device to accurately measure and monitor the discharge

from the pumping well.

- 6. Provide sufficient pipe to transport the discharge from the pumping well to an area beyond the expected cone of depression. Conducting a pumping test in contaminated groundwater may require treatment, special handling, or a discharge permit before the water can be discharged.
- 7. The discharge pipe must have a gate valve to control the pumping rate.
- 8. Determine if there is an outlet near the well head for water quality determination and sampling.

7.7.3 Pre-Test Monitoring

It is desirable to monitor pretest water levels at the test site for about 1 week prior to performance of the test. This can be accomplished by using a continuous-recording device such as a Stevens recorder. This information allows the determination of the barometric efficiency of the aquifer when barometric records are available. It also helps determine if the aquifer is experiencing an increase or decrease in head with time due to recharge or pumping in the nearby area, or diurnal effects of evapotranspiration. Changes in barometric pressure are recorded during the test (preferably with an onsite barograph) in order to correct water levels for any possible fluctuations which may occur due to changing atmospheric conditions. Pretest water level trends are projected for the duration of the test. These trends and/or barometric changes are used to "correct" water levels during the test so they are representative of the hydraulic response of the aquifer due to pumping of the test well.

7.7.4 Step Test

Conduct a step test prior to initiating a long term pumping test. The purpose of a step test is to estimate the greatest flow rate that may be sustained during a long term test. The test is performed by progressively increasing the flow rate at 1 hour intervals. The generated drawdown versus time data is plotted on semilogarithmic graph paper, and the discharge rate is determined from this graph.

7.7.5 Pump Test

Time Intervals

After the pumping well has fully recovered from the step test, the long term pumping test may start. At the beginning of the test, the discharge rate should be set as quickly and accurately as possible. The water levels in the pumping well and observation wells will be recorded according to Tables 4 and 5 below.

Water Level Measurements

Water levels will be measured as specified in ERT SOP #2151, Well Level Measurement. During the early part of the test, sufficient personnel should be

available to have at least one person at each observation well and at the pumping well. After the first 2 hours, two people are usually sufficient to continue the test. It is not necessary that readings at the wells be taken simultaneously. It is very important that depth to water readings be measured accurately and readings recorded at the exact time measured. Alternately, individual pressure transducers and electronic data-loggers may be used to reduce the number of field personnel hours required to complete the pumping test. A typical aquifer pump test form is shown in Appendix C.

During a pumping test, the following data must be recorded accurately on the aquifer test data form.

1. Site ID -- A number assigned to identity a specific site.

Table 4: Time Intervals for Measuring Drawdown in the Pumped Well

Elapsed Time From Start of Test (Minutes)	Interval Between Measurements (Minutes)
0 - 10	0.5 - 1
10 - 15	1
15 - 60	5
60-300	30
300-1440	60
1440 - termination	480

Table 5: Time Intervals for Measuring Drawdown in an Observation Well

Elapsed Time From Start of Test (Minutes)	Interval Between Measurements (Minutes)
O-60	2
60 - 120	5
120 - 240	10
240 - 360	30
360-1440	60
1440 - termination	480

- 2. Location -- The location of the well in which water level measurements are being taken.
- 3. Distance from Pumped Well -- Distance between the observation well and the pumping well in feet.
- 4. Logging Company -- The company conducting the pumping test.
- 5. Test Start Date -- The date when the pumping test began.
- 6. Test Start Time -- Start time, using a 24hour clock.
- 7. Static Water Level (Test Start) -- Depth to water, in feet and tenths of feet, in the observation well at the beginning of the pumping test.
- 8. Test End Date -- The date when the pumping test was completed.
- 9. Test End Time -- End time, using a 24hour clock.
- 10. Static Water Level (Test End) -- Depth to water, in feet and tenths of feet, in the observation well at the end of the pumping test.
- 11. Average Pumping Rate -- Summation of all entries recorded in the Pumping Rate (gal/min) column divided by the total number of Pumping Rate (gal/min) readings.
- 12. Measurement Methods -- Type of instrument used to measure depth-to-water (this may include steel tape, electric sounding probes, Stevens recorders, or pressure transducers).
- 13. Comments -- Appropriate observations or information which have not been recorded elsewhere, including notes on sampling.
- 14. Elapsed Time (min) -- Time of measurement recorded continuously from start of test (time 00.00).
- 15. Depth to Water (ft) -- Depth to water, in feet and tenths of feet, in the observation well at the time of the water level measurement.
- 16. Pumping Rate (gal/min) -- Plow rate of pump measured from an orifice, weir, flow meter,

container or other type of water-measuring device.

Test Duration

The duration of the test is determined by the needs of the project and properties of the aquifer. One simple test for determining adequacy of data is when the log-time versus drawdown for the most distant observation well begins to plot as a straight line on the semilogarithmic graph paper. There are several exceptions to this simple rule of thumb, therefore, it should be considered a minimum criterion. Different hydrogeologic conditions can produce straight line trends on log-time versus drawdown plots. In general, longer tests produce more definitive results. A duration of 1 to 3 days is desirable, followed by a similar period of monitoring the recovery of the water level. Unconfined aquifers and partially penetrating wells may have shorter test durations. Knowledge of the local hydrogeology, combined with a clear understanding of the overall project objectives, is necessary in interpreting just how long the test should be conducted. There is no need to continue the test if the water level becomes constant with time. This normally indicates that a hydrogeologic source has been intercepted and that additional useful information will not be collected by continued pumping.

7.7.6 Post Operation

- 1. After completion of water level recovery measurements, decontaminate and/or dispose of equipment as per ERT SOP #2006, Sampling Equipment Decontamination.
- 2. When using an electronic data-logger, use the following procedures.
 - see Stop logging sequence.
 - Print data, or save memory and disconnect battery at the end of the day's activities.
- 3. Replace testing equipment in storage containers.
- 4. Check sampling equipment and supplies. Repair or replace all broken or damaged equipment.
- 5. Review field forms for completeness.

6. Interpret pumping/recovery test field results.

7.8 CALCULATIONS

There are several accepted methods for determining aquifer properties such as transmissivity, storativity, and conductivity. However, the method to use is dependent on the characteristics of the aquifer being tested (confined, unconfined, leaky confining layer, etc.). When reviewing pump test data, texts by Fetter, or Driscoll or Freeze and Cherry may be used to determine the method most appropriate to your case. See the reference section on page 69.

7.9 QUALITY ASSURANCE/ QUALITY CONTROL

Calibrate all gauges, transducers, flow meters, and other equipment used in conducting pumping tests before use at the site. Obtain records of the instrument calibration and file with the test data records. The calibration records will consist of laboratory measurements. If necessary, perform any on-site zero adjustment and/or calibration. Where possible, check all flow and measurement meters on-site using a container of measured volume and stopwatch; the accuracy of the meters must be verified before testing proceeds.

7.10 DATA VALIDATION

This section is not applicable to this SOP.

7.11 HEALTH AND SAFETY

8.0 SLUG TEST: SOP #2158

8.1 SCOPE AND APPLICATION

This procedure can determine the horizontal hydraulic conductivity of distinct geologic horizons under in situ conditions. The hydraulic conductivity (K) is an important parameter for modeling the flow of groundwater in an aquifer.

8.2 METHOD SUMMARY

A slug test involves the instantaneous injection of a slug (a solid cylinder of known volume) or withdrawal of a volume of water. A slug displaces a known volume of water from a well and measures the artificial fluctuation of the groundwater level.

There are several advantages to using slug tests to estimate hydraulic conductivities. First, estimates can be made in situ, thereby avoiding errors incurred in laboratory testing of disturbed soil samples. Second, compared with pump tests, slug tests can be performed quickly and at relatively low cost, because pumping and observation wells are not required. And last, the hydraulic conductivity of small discrete portions of an aquifer can be estimated (e.g., sand layers in a clay).

8.3 SAMPLE PRESERVATION, CONTAINERS, HANDLING, AND STORAGE

This section is not applicable to this Standard Operating Procedure (SOP).

8.4 INTERFERENCES AND POTENTIAL PROBLEMS

- Only the hydraulic conductivity of the area immediately surrounding the well is estimated, which may not be representative of the average hydraulic conductivity of the area.
- The storage coefficient, S, usually cannot be determined by this method.

8.5 EQUIPMENT/APPARATUS

The following equipment is needed to perform slug tests. All equipment which comes in contact with the well should be decontaminated and tested prior to commencing field activities.

- tape measure (subdivided into tenths of feet)
- water pressure transducer
- electric water level indicator
- weighted tapes
- steel tape (subdivided into tenths of feet)
- electronic data-logger (if transducer method is used)
- stainless steel slug of a known volume
- watch or stopwatch with second hand
- semilogarithmic graph paper (if required)
- waterproof ink pen and logbook
- thermometer
- appropriate references and calculator
- electrical tape
- 21X micrologger
- Compaq portable computer or equivalent with Grapher installed on the hard disk

8.6 REAGENTS

No chemical reagents are used in this procedure; however, decontamination solvents may be necessary. When decontaminating the slug or equipment, refer to ERT SOP #2006, Sampling Equipment Decontamination, and the site-specific work plan.

8.7 PROCEDURES

8.7.1 Field Procedures

When the slug test is performed using an electronic data-logger and pressure transducer, all data will be stored internally or on computer diskettes or tape. The information will be transferred directly to the main computer and analyzed. Keep a computer printout of the data in the files as documentation.

If the slug test data is collected and recorded manually, the slug test data form (Appendix C) will

be used to record observations. The slug test data form should include the following information:

- site ID -- identification number assigned to the site
- zz location ID -- identification of location being tested
- date -- the date when the test data were collected in this order: year, month, day (e.g., 900131 for January 31, 1990)
- slug volume (ft²) = manufacturer's specification for the known volume or displacement of the slug device
- logger -- identifies the company or person responsible for performing the field measurements
- test method -- the slug device either is injected or lowered into the well, or is withdrawn or pulled-out from the monitor well. Check the method that is applicable to the test situation being run.
- comments -- appropriate observations or information for which no other blanks are provided.
- elapsed time (minutes) -- cumulative time readings from beginning of test to end of test, in minutes
- depth to water (feet) -- depth to water recorded in tenths of feet

The following general procedures may be used to collect and report slug test data. These procedures may be modified to reflect site-specific conditions:

- 1. Decontaminate the transducer and cable.
- 2. Make initial water level measurements on monitoring **wells** in an upgradient-to-downgradient sequence, if possible, to minimize the potential for cross-contamination.
- 3. Before beginning the slug test, record information into the electronic data-logger. The type of information may vary depending on the model used. When using different model, consult the operator's manual for the proper data entry sequence to be used.
- 4. Test wells from least contaminated to most contaminated, if possible.
- Determine the static water level in the well by measuring the depth to water periodically for several minutes and taking the average of the readings, (see SOP #2151, Water Level

Measurement).

- 6. Cover sharp edges of the well casing with duct tape to protect the transducer cables.
- 7. Install the transducer and cable in the well to a depth below the target drawdown estimated for the test but at least 2 feet from the bottom of the well. Be sure the depth of submergence is within the design range stamped on the transducer. Temporarily tape the transducer cable to the well to keep the transducer at a constant depth.
- 8. Connect the transducer cable to the electronic data-logger.
- 9. Enter the initial water level and transducer design range into the recording device according to the manufacturer's instructions. The transducer design range will be stamped on the side of the transducer. Record the initial water level on the recording device.
- 10. "Instantaneously" introduce or remove a known volume or slug of water to the well. Another method is to introduce a solid cylinder of known volume to displace and raise the water level, allow the water level to restabilize and remove the cylinder. It is important to remove or add the volumes as quickly as possible because the analysis assumes an "instantaneous" change in volume is created in the well.
- 11. Consider the moment of volume addition or removal as time zero. Measure and record the depth to water and the time at each reading. Depths should be measured to the nearest 0.01 foot. The number of depth-time measurements necessary to complete the test is variable. It is critical to make as many measurements as possible in the early part of the test. The number and intervals between measurements will be determined from previous aquifer tests or evaluations.
- 12. Continue measuring and recording depth-time measurements until the water level returns to equilibrium conditions or a sufficient number of readings have been made to clearly show a trend on a semilogarithmic plot of time versus depth.
- 13. Retrieve slug (if applicable).

Note: The time required for a slug test to be completed is a function of the volume of the slug, the hydraulic conductivity of the formation and the type of well completion. The slug volume should be large enough that a sufficient number of water level measurements can be made before the water level returns to equilibrium conditions. The length of the test may range from less than a minute to several hours. If the well is to be used as a monitoring well, precautions against contaminating it should be taken. If water is added to the monitoring well, it should be from an uncontaminated source and transported in a clean container. Bailers or measuring devices should be decontaminated prior to the test. If tests are performed on more than one monitoring well, care must be taken to avoid cross-contamination of the wells.

Slug tests should be conducted on relatively undisturbed wells. If a test is conducted on a well that has recently been pumped for water sampling purposes, the measured water level must be within 0.1 foot of the static water level prior to sampling. At least 1 week should elapse between the drilling of a well and the performance of a slug test.

8.7.2 Post Operation

When using an electronic data-logger, use the following procedure:

- 1. Stop logging sequence.
- 2. Print data.
- 3. Send data to computer by telephone.
- 4. Save memory and disconnect battery at the end of the day's activities.
- 5. Review field forms for completeness.

8.8 CALCULATIONS

The simplest interpretation of piezometer recovery is that of Hvorslev (1951). The analysis assumes a homogenous, isotropic medium in which soil and water are incompressible. Hvorslev's expression for hydraulic conductivity (K) is:

$$K=r21n(L/R)$$

 $2LT_{\circ}$

for L/R > 8

where:

K = hydraulic conductivity [feet/second]

= casing radius [feet]

r = length of open screen (or open borehole)
[feet]

R = filter pack (borehole) radius [feet]

 T_{\circ} = Basic Time Lag [seconds]; value of t on semilogarithmic plot of (H-h)/(H-H $_{\circ}$) vs. t, when (H-h)/(H-H $_{\circ}$) = 0.37

where:

H = initial water level prior to removal of slug

 H_{\circ} = water level at t = 0

h = recorded water level at t> 0

(Hvorslev, 1951; Freeze and Cherry, 1979)

The Bower and Rice method is also commonly used for K calculations. However, it is much more time consuming than the Hvorslev method. Refer to Freeze and Cherry or Fetter for a discussion of these methods.

8.9 QUALITY ASSURANCE/ QUALITY CONTROL

The following general quality assurance procedures apply:

- All data must be documented on standard chain of custody forms, field data sheets, or within personal/site logbooks.
- All instrumentation must be operated in accordance with operating instructions as supplied by the manufacturer, unless otherwise specified in the work plan. Equipment checkout and calibration activities must occur prior to sampling/operation, and they must be documented.

The following specific quality assurance activity will apply:

Each well should be tested at least twice in order to compare results.

8.10 DATA VALIDATION

This section is not applicable to this SOP.

8.11 HEALTH AND SAFETY

APPENDIX A

Sampling Train Schematic

Figure 1: Sampling Train Schematic SOP #2149

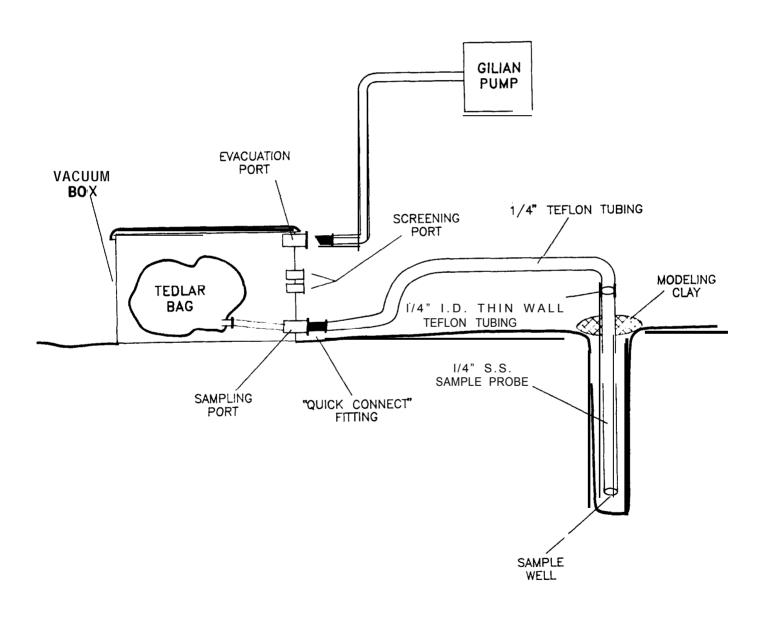
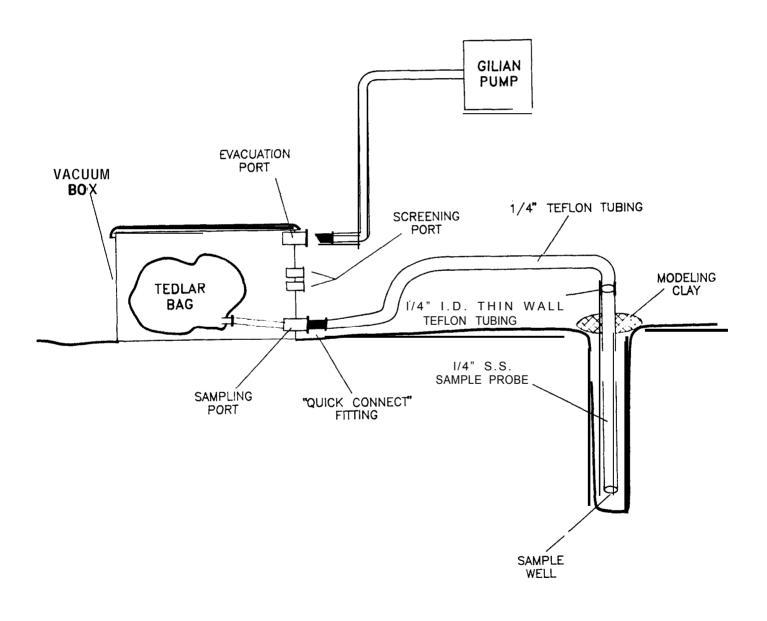


Figure 1: Sampling Train Schematic SOP #2149



APPENDIX A

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[feet]

R = filter pack (borehole) radius [feet]

 T_{\circ} = Basic Time Lag [seconds]; value of t on semilogarithmic plot of (H-h)/(H-H $_{\circ}$) vs. t, when (H-h)/(H-H $_{\circ}$) = 0.37

where:

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(Hvorslev, 1951; Freeze and Cherry, 1979)

The Bower and Rice method is also commonly used for K calculations. However, it is much more time consuming than the Hvorslev method. Refer to Freeze and Cherry or Fetter for a discussion of these methods.

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Each well should be tested at least twice in order to compare results.

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- logger -- identifies the company or person responsible for performing the field measurements
- test method -- the slug device either is injected or lowered into the well, or is withdrawn or pulled-out from the monitor well. Check the method that is applicable to the test situation being run.
- comments -- appropriate observations or information for which no other blanks are provided.
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The following general procedures may be used to collect and report slug test data. These procedures may be modified to reflect site-specific conditions:

- 1. Decontaminate the transducer and cable.
- 2. Make initial water level measurements on monitoring **wells** in an upgradient-to-downgradient sequence, if possible, to minimize the potential for cross-contamination.
- 3. Before beginning the slug test, record information into the electronic data-logger. The type of information may vary depending on the model used. When using different model, consult the operator's manual for the proper data entry sequence to be used.
- 4. Test wells from least contaminated to most contaminated, if possible.
- Determine the static water level in the well by measuring the depth to water periodically for several minutes and taking the average of the readings, (see SOP #2151, Water Level

Measurement).

- 6. Cover sharp edges of the well casing with duct tape to protect the transducer cables.
- 7. Install the transducer and cable in the well to a depth below the target drawdown estimated for the test but at least 2 feet from the bottom of the well. Be sure the depth of submergence is within the design range stamped on the transducer. Temporarily tape the transducer cable to the well to keep the transducer at a constant depth.
- 8. Connect the transducer cable to the electronic data-logger.
- 9. Enter the initial water level and transducer design range into the recording device according to the manufacturer's instructions. The transducer design range will be stamped on the side of the transducer. Record the initial water level on the recording device.
- 10. "Instantaneously" introduce or remove a known volume or slug of water to the well. Another method is to introduce a solid cylinder of known volume to displace and raise the water level, allow the water level to restabilize and remove the cylinder. It is important to remove or add the volumes as quickly as possible because the analysis assumes an "instantaneous" change in volume is created in the well.
- 11. Consider the moment of volume addition or removal as time zero. Measure and record the depth to water and the time at each reading. Depths should be measured to the nearest 0.01 foot. The number of depth-time measurements necessary to complete the test is variable. It is critical to make as many measurements as possible in the early part of the test. The number and intervals between measurements will be determined from previous aquifer tests or evaluations.
- 12. Continue measuring and recording depth-time measurements until the water level returns to equilibrium conditions or a sufficient number of readings have been made to clearly show a trend on a semilogarithmic plot of time versus depth.
- 13. Retrieve slug (if applicable).

8.0 SLUG TEST: SOP #2158

8.1 SCOPE AND APPLICATION

This procedure can determine the horizontal hydraulic conductivity of distinct geologic horizons under in situ conditions. The hydraulic conductivity (K) is an important parameter for modeling the flow of groundwater in an aquifer.

8.2 METHOD SUMMARY

A slug test involves the instantaneous injection of a slug (a solid cylinder of known volume) or withdrawal of a volume of water. A slug displaces a known volume of water from a well and measures the artificial fluctuation of the groundwater level.

There are several advantages to using slug tests to estimate hydraulic conductivities. First, estimates can be made in situ, thereby avoiding errors incurred in laboratory testing of disturbed soil samples. Second, compared with pump tests, slug tests can be performed quickly and at relatively low cost, because pumping and observation wells are not required. And last, the hydraulic conductivity of small discrete portions of an aquifer can be estimated (e.g., sand layers in a clay).

8.3 SAMPLE PRESERVATION, CONTAINERS, HANDLING, AND STORAGE

This section is not applicable to this Standard Operating Procedure (SOP).

8.4 INTERFERENCES AND POTENTIAL PROBLEMS

- Only the hydraulic conductivity of the area immediately surrounding the well is estimated, which may not be representative of the average hydraulic conductivity of the area.
- The storage coefficient, S, usually cannot be determined by this method.

8.5 EQUIPMENT/APPARATUS

The following equipment is needed to perform slug tests. All equipment which comes in contact with the well should be decontaminated and tested prior to commencing field activities.

- tape measure (subdivided into tenths of feet)
- water pressure transducer
- electric water level indicator
- weighted tapes
- steel tape (subdivided into tenths of feet)
- electronic data-logger (if transducer method is used)
- stainless steel slug of a known volume
- watch or stopwatch with second hand
- semilogarithmic graph paper (if required)
- waterproof ink pen and logbook
- thermometer
- appropriate references and calculator
- electrical tape
- 21X micrologger
- Compaq portable computer or equivalent with Grapher installed on the hard disk

8.6 REAGENTS

No chemical reagents are used in this procedure; however, decontamination solvents may be necessary. When decontaminating the slug or equipment, refer to ERT SOP #2006, Sampling Equipment Decontamination, and the site-specific work plan.

8.7 PROCEDURES

8.7.1 Field Procedures

When the slug test is performed using an electronic data-logger and pressure transducer, all data will be stored internally or on computer diskettes or tape. The information will be transferred directly to the main computer and analyzed. Keep a computer printout of the data in the files as documentation.

If the slug test data is collected and recorded manually, the slug test data form (Appendix C) will

6. Interpret pumping/recovery test field results.

7.8 CALCULATIONS

There are several accepted methods for determining aquifer properties such as transmissivity, storativity, and conductivity. However, the method to use is dependent on the characteristics of the aquifer being tested (confined, unconfined, leaky confining layer, etc.). When reviewing pump test data, texts by Fetter, or Driscoll or Freeze and Cherry may be used to determine the method most appropriate to your case. See the reference section on page 69.

7.9 QUALITY ASSURANCE/ QUALITY CONTROL

Calibrate all gauges, transducers, flow meters, and other equipment used in conducting pumping tests before use at the site. Obtain records of the instrument calibration and file with the test data records. The calibration records will consist of laboratory measurements. If necessary, perform any on-site zero adjustment and/or calibration. Where possible, check all flow and measurement meters on-site using a container of measured volume and stopwatch; the accuracy of the meters must be verified before testing proceeds.

7.10 DATA VALIDATION

This section is not applicable to this SOP.

7.11 HEALTH AND SAFETY

- 2. Location -- The location of the well in which water level measurements are being taken.
- 3. Distance from Pumped Well -- Distance between the observation well and the pumping well in feet.
- 4. Logging Company -- The company conducting the pumping test.
- 5. Test Start Date -- The date when the pumping test began.
- 6. Test Start Time -- Start time, using a 24hour clock.
- 7. Static Water Level (Test Start) -- Depth to water, in feet and tenths of feet, in the observation well at the beginning of the pumping test.
- 8. Test End Date -- The date when the pumping test was completed.
- 9. Test End Time -- End time, using a 24hour clock.
- 10. Static Water Level (Test End) -- Depth to water, in feet and tenths of feet, in the observation well at the end of the pumping test.
- 11. Average Pumping Rate -- Summation of all entries recorded in the Pumping Rate (gal/min) column divided by the total number of Pumping Rate (gal/min) readings.
- 12. Measurement Methods -- Type of instrument used to measure depth-to-water (this may include steel tape, electric sounding probes, Stevens recorders, or pressure transducers).
- 13. Comments -- Appropriate observations or information which have not been recorded elsewhere, including notes on sampling.
- 14. Elapsed Time (min) -- Time of measurement recorded continuously from start of test (time 00.00).
- 15. Depth to Water (ft) -- Depth to water, in feet and tenths of feet, in the observation well at the time of the water level measurement.
- 16. Pumping Rate (gal/min) -- Plow rate of pump measured from an orifice, weir, flow meter,

container or other type of water-measuring device.

Test Duration

The duration of the test is determined by the needs of the project and properties of the aquifer. One simple test for determining adequacy of data is when the log-time versus drawdown for the most distant observation well begins to plot as a straight line on the semilogarithmic graph paper. There are several exceptions to this simple rule of thumb, therefore, it should be considered a minimum criterion. Different hydrogeologic conditions can produce straight line trends on log-time versus drawdown plots. In general, longer tests produce more definitive results. A duration of 1 to 3 days is desirable, followed by a similar period of monitoring the recovery of the water level. Unconfined aquifers and partially penetrating wells may have shorter test durations. Knowledge of the local hydrogeology, combined with a clear understanding of the overall project objectives, is necessary in interpreting just how long the test should be conducted. There is no need to continue the test if the water level becomes constant with time. This normally indicates that a hydrogeologic source has been intercepted and that additional useful information will not be collected by continued pumping.

7.7.6 Post Operation

- 1. After completion of water level recovery measurements, decontaminate and/or dispose of equipment as per ERT SOP #2006, Sampling Equipment Decontamination.
- 2. When using an electronic data-logger, use the following procedures.
 - see Stop logging sequence.
 - Print data, or save memory and disconnect battery at the end of the day's activities.
- 3. Replace testing equipment in storage containers.
- 4. Check sampling equipment and supplies. Repair or replace all broken or damaged equipment.
- 5. Review field forms for completeness.

APPENDIX B

HNU Field Protocol

HNU Field Protocol SOP #2149

Startup Procedure

- Before attaching the probe, check the function switch on the control panel to ensure that it is in the 'off position. Attach the probe by plugging it into the interface on the top of the readout module. Use care in aligning the prongs in the probe cord with the plug in; do not force.
- Turn the function switch to the battery check position. The needle on the meter should read within or above the green area on the scale. If not, recharge the battery. If the red indicator light comes on, the battery needs recharging.
- 3. Turn the function switch to any range setting. For no more than 2 to 3 seconds look into the end of the probe to see if the lamp is on. If it is on, you will see a purple glow. Do not stare into the probe any longer than three seconds. Long term exposure to UV light can damage eyes. Also, listen for the hum of the fan motor.
- 4. To zero the instrument, turn the function switch to the standby position and rotate the zero adjustment until the meter reads zero. A calibration gas is not needed since this is an electronic zero adjustment. If the span adjustment setting is changed after the zero is set, the zero should be rechecked and adjusted, if necessary. Wait 15 to 20 seconds to ensure that the zero reading is stable. If necessary, readjust the zero.

Operational Check

- 1. Follow the startup procedure.
- 2. With the instrument set on the 0-20 range, hold a solvent-based Magic Marker near the probe tip. If the meter deflects upscale, the instrument is working.

Field Calibration Procedure

1. Follow the startup procedure and the operational check.

- 2. Set the function switch to the range setting for the concentration of the calibration gas.
- 3. Attach a regulator (HNU 101-351) to a disposable cylinder of isobutylene gas. Connect the regulator to the probe of the HNU with a piece of clean Tygon tubing. Turn the valve on the regulator to the 'on' position.
- 4. After 15 seconds, adjust the span dial until the meter reading equals the concentration of the calibration gas used. The calibration gas is usually 100 ppm of isobutylene in zero air. The cylinders are marked in benzene equivalents for the 10.2 eV probe (approximately 55 ppm benzene equivalent) and for the 11.7 eV probe (approximately 65 ppm benzene equivalent). Be careful to unlock the span dial before adjusting it. If the span has to be set below 3.0 calibration, the lamp and ion chamber should be inspected and cleaned as appropriate. For cleaning of the 11.7 eV probe, only use an electronic-grade, oil-free freon or similar waterfree, grease-free solvent.
- 5. Record in the field log: the instrument ID # (EPA decal or serial number if the instrument is a rental); the initial and final span settings; the date and time; concentration and type of calibration used; and the name of the person who calibrated the instrument.

Operation

- 1. Follow the startup procedure, operational check, and calibration check.
- Set the function switch to the appropriate range. If the concentration of gases or vapors is unknown, set the function switch to the 0-20 ppm range. Adjust it if necessary.
- 3. While taking care not to permit the HNU to be exposed to excessive moisture, dirt, or contamination, monitor the work activity as specified in the site health and safety plan.
- When the activity is completed or at the end of the day, carefully clean the outside of the HNU with a damp disposable towel to remove any

- visible dirt. Return the HNU to a secure area and place on charge.
- 5. With the exception of the probe's inlet and exhaust, the HNU can be wrapped in clear

plastic to prevent it from becoming contaminated and to prevent water from getting inside in the event of precipitation.

APPENDIX C

Forms

Well Completion Form

SOP #2150

ć.				L INSTALL		
Clienti						
Total	Depth	·	Casing Size & T	ype:	s c	reen <u>Size</u>
Conne	nts:	·				
	>				Comple	tion Data
Depth	Symbol Stratigraphy	Sample	Description			
ſ						

SITE NAME:					
LOG DATE:			LOGG	ER NAME:	
MEASUREMENT REFERENCE POINT:		-TO	-TOP OF GROUND		-TOP OF CASING
LOCATION	TIME	DEPT WATE			COMMENTS
			· · · · · · · · · · · · · · · · · · ·		

Pump/Recovery Test Data Sheet

SOP #2157

SITE ID:		DISTANCE FROM PUMPED WELL (FT):					
LOCATION:			LOGGER:				
TEST START			TEST EN	ND			
DATE:			DAT	E :			
TIME:		I	TIM	E:			
STATIC WATER	LEVEL (FT):		STA'	TIC WATER LEVEL	(FT):		
AVERAGE PUMI	PING RATE (GAL/M	IIN):					
MEASUREMENT	METHODS:						
COMMENTS:							
ELAPSED TIME	PUMP TEST DEPTH TO		PING ATE	RECOVERY TEST ELAPSED	ДЕРТН ТО		
(MIN)	WATER (FT)		/MIN)	TIME (MIN)	WATER (FT)		
0.00				0.00			

Pump/Recovery Test Data Sheet (Continued)

SOP #2157

SITE ID:			DATE:			
LOCATION:			LOGGER:			
ELAPSED TIME (MIN)	PUMP TEST DEPTH TO WATER (FT)	PUMPING RATE (GAL/MIN)		RECOVERY TEST ELAPSED TIME (MIN)	DEPTH TO WATER (FT)	
:						

Slug Test Data Form

SOP #2158

DATE:		Ī		
SITE ID:		SLUG VOLUME (FT³):		
LOCATION ID:		LOGGER:		
TEST METHOD:	_ SLUG INJECTI	ON _ SLUG WI	ГHDRAWAL	
COMMENTS:				
TIME (Begin Test #1):		TIME (Begin Test #2):		
TIME (End Test #1):		TIME (End Test #2):		
ELAPSED TIME (MIN)	DEPTH TO WATER (FT)	ELAPSED TIME (MIN)	DEPTH TO WATER (FT)	

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APPENDIX D LABORATORY ANALYTE LIST AND REPORTING LIMITS

Summit Environmental Technologies MDL Data

MDL's listed are for water, soil, sediment and biota

	sted are for war			
Analyte	Method	MDL	Units	mg/Kg
Cd	6010/200.7	0.82	ug/L	0.02
Cr	6010/200.7	1.09	ug/L	0.05
Ва	6010/200.7	0.57	ug/L	0.03
As	6010/200.7	2.06	ug/L	0.19
Cu	6010/200.7	1.83	ug/L	0.08
Zn	6010/200.7	0.53	ug/L	0.08
Ni	6010/200.7	1.16	ug/L	0.07
TI	6010/200.7	5.29	ug/L	0.16
Ве	6010/200.7	0.72	ug/L	0.08
Pb	6010/200.7	2.1	ug/L	0.45
Se	6010/200.7	4.49	ug/L	0.1
Sb	6010/200.7	2.37	ug/L	0.08
Ca	6010/200.7	3.93	ug/L	0.68
Mg	6010/200.7	0.79	ug/L	0.52
Mn	6010/200.7	0.43	ug/L	0.06
Co	6010/200.7	0.51	ug/L	0.06
Fe	6010/200.7	10.56	ug/L	0.64
Al	6010/200.7	10.53	ug/L	4.38
V	6010/200.7	1.74	ug/L	0.05
Mo	6010/200.7	2.87	ug/L	0.1
Ti	6010/200.7	1.49	ug/L	0.06
Ag	6010/200.7	0.39	ug/L	0.04
В	6010/200.7	0.82	ug/L	0.31
K	6010/200.7	23.99	ug/L	7.24
Na	6010/200.7	197.41	ug/L	43.49
Si	6010/200.7	2.97	ug/L	1.53
Sn	6010/200.7	37.69	ug/L	0.87
Р	6010/200.7	17.35	ug/L	1.1
Analyte	Method	MDL	Units	
Low Level Hg	1631	0.5	ng/L	N/A
Analyte	Method	MDL	Units	
Fluoride	300.0/9056	0.08	mg/L	0.08
Chloride	300.0/9056	0.03	mg/L	0.03
Nitrite	300.0/9056	0.03	mg/L	0.03
Bromide	300.0/9056	0.04	mg/L	0.04
Nitrate	300.0/9056	0.03	mg/L	0.03
Phosphate	300.0/9056	0.11	mg/L	0.11
Sulfate	300.0/9056	0.09	mg/L	0.07

Analysis	Water MDC (pCi/L)	Solid MDC (pCi/g)
Isotopic Uranium		
U-234	0.06	0.1
U-235	0.06	0.1
U-236	0.06	0.1
U-238	0. 06	0.1
Gamma Spectroscopy (example isotopes)		
Ba-133	4.0	0.12
Bi-211	14.2	0.08
Bi-212	52.4	0.19
Bi-214	63.5	0.18
Cd-109	46.2	0.24
Co-57	6.9	0.04
Co-60	3.5	0.01
Cs-134	4.1	0.02
Cs-137	3.7	0.02
Eu-152	9.7	0.04
Eu-154	4.1	0.02
Eu-155	6.2	0.03
I-129	6.6	0.04
I-131	4.8	0.10
K-40	23.7	0.09
Mn-54	4.1	0.01
Na-22	3.8	0.02
Pb-210	55.9	0.23
Pb-212	4.2	0.02
Pb-214	4.7	0.03
Ra-226	48.3	0.3
Ra-228	11.5	0.01
Sc-46	3.8	0.02
Zn-65	9.5	0.09

APPENDIX E

START-3 TDD No. TO-0019-10-11-01



START3 Technical Direction Document

TDD #: TO-0019-10-11-01 Contract: EP-W-06-042

Assessment/Inspection Activities (CERCLA PIPELINE FUNDS - FOR SITE ASSESSMENT GROUP USE) (0019) Weston Solutions, Inc.

	Į	=	req	uired	field
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TDD Name:	Jackpile-Paguate Urar ESI	nium Mine	! Period:	Base Period	
! Purpose:	Work Assignment Initia	ation			
! Priority:	High		! Start Date:	11/29/2010	
Overtime:	Yes		! Completion Date:	08/01/2011	
! Funding Category:	Removal		Invoice Unit:		
! Project/Site Name:	Jackpile-Paguate Urar	nium Mine	WorkArea:	ASSESSMENT/INSPECTION ACTIVITIES	ONS
Project Address:	SR 279, Near Paguate Pueblo	e, Laguna	Activity:	Expanded Site Inspection (E	ESI)
County:	Cibola		Work Area Code:		
City, State:	Paguate; NM,		Activity Code:	ES	
Zip:			EMERGENCY CODE:	\square kat \square rit	
! SSID:	A6T3		FPN:		
CERCLIS:	NMN000607033		Performance Based:	No	
Operable Unit:					
Authorized TDD Ce	eiling:		Cost/	Fee	LOE (Hours)
	Previous Action(s):		\$0	0.00	0.0
	This Action:		\$100,000	0.00	0.0
	New Total:		\$100,000	0.00	0.0

Specific Elements Perform ESI activities in accordance with EPA 540-R092-021., Further characterize and define a site for potential response action., Obtain additional information required to support preparation of HRS package for NPL listing which requires scoring of the site.

Description of Work:

All activities performed in support of this TDD shall be in accordance with the contract and TO PWS.

The contractor shall develop a work plan/cost estimate for completing a ESI at the Jackpile Mine in Paquate, New Mexico. Specific tasks in the work plan/cost estimate shall include:

- Management/Cost Estimate;
- 2. QASP/HASP/Lab Procurement;
- 3. Field Activities
- 4. ESI Report
- 5. HRS documentation package

Prior to developing workplan/cost estimate the contractor shall review all existing information (including review of information used in the draft HRS package and SI report) and attend scoping meeting with EPA project team. In addition to developing a work plan/cost estimate, the contractor shall develop a Quality Assurance Sampling Plan (QASP) and Health and Safety Plan (HASP) and procure a laboratory to conduct chemical analyses prior to initiating field activities at the site. The laboratory shall provide a defensible analytical data package(s) for any samples collected during the

execution of the field activities under this TDD.

Field activities shall focus on the collection and analysis ground water and surface water samples at the site to aid in the development of a conceptual site model for the ground water to surface water migration and an understanding of background conditions at the mine. The contractor shall ensure that a person/persons with hydrology and mining experience and geochemistry be part of the site team. It is estimated that no more than 100 samples (including QA/QC samples) will be collected during the field activities. Samples shall be transported/shipped to the procured laboratory for both metals, elemental uranium and activity. A 2-week turnaround time shall be requested for the analytical results. Field activities shall include a limited survey of the area surrounding the site in order to verify potential pathway targets and off-site conditions.

An draft ESI report shall be developed for the site. Analytical data collected under this TDD shall be validated and defensible data validation reports developed. The report shall incorporate the validated analytical results from the samples collected under this TDD, as well as analytical results from samples collected as part of the SI. The report shall be developed according to the EPA guidance for performing site inspections (EPA540-R-92-021, Directive 9345.1-05). A draft report for the site shall be submitted to EPA for review no later than June 13, 2011.

All activities shall be coordinated with the EPA WAM, Brenda Cook (214-665-7436). The START contractor shall contact Brenda Cook after receipt of this TDD.

Accounting and Appropriation Information

SFO: 22

-											51 01 22
	Line	DCN	IFMS	Budget/ FY	Appropriati on Code	Budget Org Code	Program Element	Object Class	Site Project	Cost Org Code	Amount
	2	PLC036 PLC026 PLC016	XXX XXX XXX	10 09 10	Т	6A00P	302DD2C 302DD2C 302DD2C	2505	A6T3ES00	C001 C001 C001	\$65,000.00 \$1,700.00
ı	3	Leoro	ΛΛΛ	10	Licb	07 1001	3020020	2505	10131500	2001	\$33,300.00

Funding Summary:	Funding
Previous:	\$0.00
This Action:	\$100,000.00
Total:	\$100,000.00

Funding Category Removal

Section

- Signed by Brenda Cook/R6/USEPA/US on 11/23/2010 09:14:20 AM, according to Cheng Wei Feng/start

Date: 11/23/2010 : Brenda Cook

Project Officer Section - Signed by Cora Stanley/R6/USEPA/US on 11/29/2010 08:49:18 AM, according to C Project Officer: Linda Carter Date: 11/23/2010

Contracting Officer Section - Signed by Cora Stanley/R6/USEPA/US on 11/29/2010 08:49:18 AM, according

Contracting Officer: Cora Stanley Date: 11/29/2010

Contractor Section - Signed by Robert Beck/start6/rfw-start/us on 11/29/2010 11:07:10 AM, according to During the past three (3) calendar years has your company, or any of your employees that will No

be working at this site, previously performed work at this site /facility? O Yes

Contractor Contact: Robert Beck Date: 11/29/2010

APPENDIX D LABORATORY DATA PACKAGES



LABORATORY REPORT

Client

Weston Solutions 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

> Order Number 1107985 FINAL REPORT

Project Number Jackpile

Issued Monday, June 06, 2011

Total Number of Pages

80 (excluding C.O.C. and cooler receipt form)

Approved By: What

QA Manager

NELAC Accreditation #E87688



Sample Summary

Client: Weston Solutions Order Number: <u>1107985</u>

Laboratory ID	Client ID	Matrix	Sampling Date
1107985-01	JPOP41S-110421	Liquid	4/21/2011
1107985-02	MD-SW-110420	Liquid	4/20/2011
1107985-03	MW-1-110419	Liquid	4/19/2011
1107985-04	MW-2-110421	Liquid	4/21/2011
1107985-05	MW-2-2-110421	Liquid	4/21/2011
1107985-06	MW-3-110420	Liquid	4/20/2011
1107985-07	MW-4-110420	Liquid	4/20/2011
1107985-08	MW-6-110420	Liquid	4/20/2011
1107985-09	MW-7-110419	Liquid	4/19/2011
1107985-10	MW-8-110419	Liquid	4/19/2011
1107985-11	MW-MD-02-110420	Liquid	4/20/2011
1107985-12	MW-MD-110420	Liquid	4/20/2011
1107985-13	NPOP20E-110420	Liquid	4/20/2011
1107985-14	PR-SED-01-110420	Solid	4/20/2011
1107985-15	PR-SW-01-110420	Liquid	4/20/2011
1107985-16	RB-01-110420	Liquid	4/20/2011
1107985-17	RB-02-110421	Liquid	4/21/2011
1107985-18	RM-JM-SED-02-	Solid	4/20/2011
1107985-19	RM-JM-SED-110420	Solid	4/20/2011
1107985-20	RM-JM-SW-02	Liquid	4/20/2011
1107985-21	RM-JM-SW-110420	Liquid	4/20/2011
1107985-22	RM-SED-BG-110419	Solid	4/19/2011
1107985-23	RM-SW-BG-110419	Liquid	4/19/2011
1107985-24	RP-JM-SED-01	Solid	4/20/2011
1107985-25	RP-JM-SED-110420	Solid	4/20/2011
1107985-26	RP-JM-SW-01-110420	Liquid	4/20/2011
1107985-27	RP-JM-SW-110420	Liquid	4/20/2011
1107985-28	RP-SED-01-110419	Solid	4/19/2011
1107985-29	RP-SED-02-110420	Solid	4/20/2011
1107985-30	RP-SED-03-110420	Solid	4/20/2011
1107985-31	RP-SED-BG-041911	Solid	4/19/2011
1107985-32	RP-SW-01-110419	Liquid	4/19/2011
1107985-33	RP-SW-BG-041911	Liquid	4/19/2011
1107985-34	SPOP35-110421	Liquid	4/21/2011
1107985-35	MW-RM-110420	Liquid	4/21/2011



Report Narrative

Client: Weston Solutions
Order Number: 1107985

Solid sample results are reported on a DRY weight basis except as noted. No problems were encountered during analysis of this order number, except as noted.

This is a final report.

Listed below are the differences between the preliminary and final results:

Calcium was changed in samples listed below:

1107985-03 1107985-09 1107895-10 1107985-16 1107985-17

Sodium was changed in samples listed below:

1107985-06 1107985-16 1107985-17 1107985-19 1107985-22 1107985-24 1107985-25 1107985-28 1107985-29 1107985-30 1107985-31 1107985-33

Magnesium was changed in samples listed below: 1107985-06 1107985-28

Iron was changed in sample 1107985-14.

Data Qualifiers:

B = Analyte found in the method blank

J = Estimated concentration of analyte between MDL (LOD) and Reporting Limit (LOQ)

C = Analyte has been confirmed by another instrument or method E = Analyte exceeds the upper limit of the calibration curve.

D = Sample or extract was analyzed at a higher dilution

X = User defined data qualifier.

S = Surrogate out of control limits

U = Undetected

a = Not Accredited by NELAC

ND = Non Detected at LOQ

DF = Dilution Factor

Limit Of Quantitation (LOQ) = Laboratory Reporting Limit (not adjusted for dilution factor)
Limit Of Detection (LOD) = Laboratory Detection Limit

Estimated uncertainty values are available upon request.

Matrices:

A = Air C = Cream

DW = Drinking Water

L = Liquid O = Oil

O = Oil SL = Sludge

SO = Soil

S = Solid T = Tablet

TC = TCLP Extract WW = Waste Water

W = Wipe

The test results meet the requirements of the NELAC standard, except where noted. The information contained in this analytical report is the sole property of Summit Environmental Technologies, Inc. and that of the client. It cannot be reproduced in any form without the consent of Summit Environmental Technologies, Inc. or the client for which this report was issued. The results contained in this report are only representative of the samples received. Conditions can vary at different times and at different sampling conditions. Summit Environmental Technologies, Inc. is not responsible for use or interpretation of the data included herein.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

JPOP41S-110421	<u>Client ID#</u> JPOP41S-110421																				
1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	1107985-01	<u>Lab ID#</u> 1107985-01
21-Apr-11	Collected 21-Apr-11																				
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
<u>د</u>	_	\subset	D				<u>_</u>		D		U,D	C			D	د		<u>_</u>	C	C	Flag
0.0155	<0.00174	<0.00529	561	0.321	0.104	52.0	0.0086	2.84	581	0.341	<0.0106	<0.00183	0.0788	0.365	744	0.0045	0.0395	0.008	<0.00206	<0.00237	Result 5.31
mg/L																			٥.	37	
J/L	mg/L	5 mg/L	87 mg/L	<u>Units</u> mg/L																	
]/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L		mg/L L		•	Units Matrix mg/L L					
g/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7		mg/L L 200.7		•	102					
L 200.7 1	Г	Г	-	_	_	_	Г	_	Г	г	—	mg/L L	_	Г	_	L 200.7 1	L 200.7 1	L 200.7 1	mg/L L 200.7 1	mg/L L	<u>Matrix</u> L
	L 200.7 1	L 200.7 1	L 200.7 1	. L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	mg/L L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	- -	Г	_	mg/L L	mg/L L 200.7 1	<u>Matrix</u> <u>Method</u> <u>DF</u> L 200.7 1

Fax: 330-253-4489

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

MD-SW-110420 110	MD-SW-110420 110	MD-SW-110420 110	MD-SW-110420 11																		
1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02	1107985-02
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	Aluminum
C	C	ل	D	ے			د		D	د	C	C	ے	\subset	D	C	د		ے	٦	ر _
<0.00053	<0.00174	0.0062	1290	0.0019	0.0322	26.8	0.0036	0.151	198	0.0026	<0.0106	<0.00183	0.0018	<0.00109	270	<0.00082	0.0008	0.0722	0.0028	0.0079	0.0129
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L
_	_	_	_	-	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	-
200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7
	<u>—</u>		_		<u></u>	<u> </u>		<u> </u>	<u></u>	\vdash	<u> </u>	Н	_	-	_	 	<u> </u>	<u> </u>	 		-
0.00053	0.00174	0.00529	0.197	0.00039	0.00449	0.0240	0.00116	0.00043	0.00079	0.00210	0.0106	0.00183	0.00051	0.00109	0.00393	0.00082	0.00072	0.00057	0.00206	0.00237	0.0105
0.0200	0.0200	0.0200	2.00	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500	0.00500	0.00400	0.0400	0.0200	0.0200	0.100
								29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	11-May-11 VVK										

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

29-Apr-11 VVK	0.0200	0.00053	1 0	200.7	_	mg/L	<0.00053	C	Zinc	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00174	1 0	200.7	_	mg/L	<0.00174	C	Vanadium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00529	1 0.	200.7	_	mg/L	0.0212		Thallium	19-Apr-11	1107985-03	MW-1-110419
11-May-11 VVK	2.00	0.197	<u> </u>	200.7	_	mg/L	867	D	Sodium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00039	1 0	200.7	_	mg/L	0.0014	حـ	Silver	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00449	1 0	200.7	_	mg/L	0.0309		Selenium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.500	0.0240	1 (200.7	_	mg/L	7.031		Potassium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00116	1 0	200.7	_	mg/L	0.0025	د	Nickel	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00043	1 0	200.7	_	mg/L	<0.00043	_	Manganese	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.100	0.00079	1 0	200.7	_	mg/L	85.54		Magnesium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0400	0.00210	1 0	200.7	_	mg/L	<0.00210	_	Lead	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.100	0.0106	1 (200.7		mg/L	<0.0106	C	Iron	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00183	1 0	200.7	٢	mg/L	<0.00183	C	Copper	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00051	1 0	200.7	Г	mg/L	0.0022	د	Cobalt	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00109	1 0	200.7	_	mg/L	<0.00109	_	Chromium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.500	0.00393	1 0	200.7	_	mg/L	28.7	D	Calcium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.00082 0.00500	1.00082	1 0	200.7	Г	mg/L	<0.00082	_	Cadmium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.00400	0.00072	1 0	200.7	_	mg/L	<0.00072	_	Beryllium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0400	0.00057	1 0	200.7	_	mg/L	0.0180	٦	Barium	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00206	1 0	200.7	Г	mg/L	<0.00206	C	Arsenic	19-Apr-11	1107985-03	MW-1-110419
29-Apr-11 VVK	0.0200	0.00237	1 C	200.7	_	mg/L	0.0075	ر	Antimony	19-Apr-11	1107985-03	MW-1-110419
Run Analyst 29-Apr-11 VVK	<u>LOQ</u>	<u>LOD</u> 0.0105	<u>DF</u>	<u>Method</u> 200.7	Matrix L	Units mg/L	<u>Result</u> 0.0240	Flag J	<u>Analyte</u> Aluminum	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-03	<u>Client ID#</u> MW-1-110419

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29-Apr-11 VVK	0.0200	0.00053	1	200.7	_	mg/L	0.0017	د ـ	Zinc	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.0200	0.00174	1	200.7	_	mg/L	<0.00174	C	Vanadium	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.0200	0.00529	<u></u>	200.7	_	mg/L	0.0106	۲	Thallium	21-Apr-11	1107985-04	MW-2-110421
11-May-11 VVK	2.00	0.197	<u> </u>	200.7	_	mg/L	648	D	Sodium	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.0200	0.00039	1	200.7	Г	mg/L	0.0020	Z,Z	Silver	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 WK	0.0200	0.00449	1	200.7	_	mg/L	0.0317		Selenium	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 WK	0.500	0.0240	<u></u>	200.7	_	mg/L	18.7	z	Potassium	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 WK	0.0200	0.00116	<u>-</u>	200.7	_	mg/L	0.0048	ل	Nickel	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.0200	0.00043	<u></u>	200.7	_	mg/L	0.275		Manganese	21-Apr-11	1107985-04	MW-2-110421
11-May-11 VVK	0.100	0.00079	<u> </u>	200.7	_	mg/L	279	D	Magnesium	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 WK	0.0400	0.00210	⊢	200.7	Г	mg/L	0.0030	ے	Lead	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.100	0.0106	<u> </u>	200.7	_	mg/L	1.15		Iron	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 WK	0.0200	0.00183		200.7	_	mg/L	<0.00183	C	Copper	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.0200	0.00051		200.7	_	mg/L	0.0006	ل	Cobalt	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.0200	0.00109	 	200.7	_	mg/L	<0.00109	C	Chromium	21-Apr-11	1107985-04	MW-2-110421
11-May-11 VVK	0.500	0.00393	<u> </u>	200.7	_	mg/L	247	D	Calcium	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.00500	0.00082	<u> </u>	200.7	_	mg/L	<0.00082	C	Cadmium	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.00400	0.00072	_	200.7	_	mg/L	<0.00072	C	Beryllium	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.0400	0.00057	<u> </u>	200.7	_	mg/L	0.0207	ر	Barium	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.0200	0.00206	1	200.7	_	mg/L	0.0033	<i>د</i>	Arsenic	21-Apr-11	1107985-04	MW-2-110421
29-Apr-11 VVK	0.0200	0.00237	Н	200.7	_	mg/L	0.0061	ے	Antimony	21-Apr-11	1107985-04	MW-2-110421
Run Analyst 29-Apr-11 VVK	0.100	L <u>OD</u> 0.0105	DF 1	Method 200.7	Matrix L	<u>Units</u> mg/L	<u>Result</u> 0.0163	<u>Flag</u> J	<u>Analyte</u> Aluminum	Collected 21-Apr-11	<u>Lab ID#</u> 1107985-04	<u>Client ID#</u> MW-2-110421

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29-Apr-11 VVK	0.0200	0.00053	<u> </u>	200.7	_	mg/L	<0.00053	_	Zinc	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 WK	0.0200	0.00174	_	200.7	٦	mg/L	<0.00174	C	Vanadium	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0200	0.00529	<u> </u>	200.7	_	mg/L	0.0077	ے	Thallium	21-Apr-11	1107985-05	MW-2-2-110421
11-May-11 VVK	2.00	0.197	<u> </u>	200.7	_	mg/L	694	D	Sodium	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0200	0.00039	⊨	200.7	_	mg/L	<0.00039	C	Silver	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0200	0.00449	_	200.7	_	mg/L	0.0327		Selenium	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 WK	0.500	0.0240	 	200.7	_	mg/L	20.32		Potassium	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0200	0.00116	<u> </u>	200.7	_	mg/L	0.0052	ل	Nickel	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0200	0.00043	<u></u>	200.7	_	mg/L	0.294		Manganese	21-Apr-11	1107985-05	MW-2-2-110421
11-May-11 VVK	0.100	0.00079		200.7	٦	mg/L	297	D	Magnesium	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0400	0.00210	<u>-</u>	200.7	_	mg/L	<0.00210	C	Lead	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.100	0.0106	—	200.7		mg/L	1.27		Iron	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0200	0.00183	<u></u> -	200.7	_	mg/L	<0.00183	C	Copper	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0200	0.00051	<u> </u>	200.7	_	mg/L	0.0006	ر	Cobalt	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0200	0.00109	—	200.7	_	mg/L	<0.00109	C	Chromium	21-Apr-11	1107985-05	MW-2-2-110421
11-May-11 VVK	0.500	0.00393	↦	200.7	_	mg/L	267	D	Calcium	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.00500	0.00082 0.00500	<u> </u>	200.7	_	mg/L	<0.00082	_	Cadmium	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.00400	0.00072		200.7	_	mg/L	<0.00072	C	Beryllium	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 VVK	0.0400	0.00057	ᆢ	200.7	_	mg/L	0.0221	د	Barium	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 WK	0.0200	0.00206	⊷	200.7	_	mg/L	<0.00206	C	Arsenic	21-Apr-11	1107985-05	MW-2-2-110421
29-Apr-11 WK	0.0200	0.00237	<u> </u>	200.7	_	mg/L	0.0057	ر	Antimony	21-Apr-11	1107985-05	MW-2-2-110421
Run Analyst 29-Apr-11 VVK	0.100 LOQ	F <u>LOD</u> 0.0105	<u>DF</u>	Method 200.7	Matrix L	Units mg/L	Result 0.0164	Flag J	<u>Analyte</u> Aluminum	Collected 21-Apr-11	<u>Lab ID#</u> 1107985-05	Client ID# MW-2-2-110421
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200.7 1 0.00053 0.0200	mg/L L	0.0066	ر ر	Zinc	20-Apr-11	1107985-06	MW-3-110420
200.7	mg/L L	<0.00174	C	Vanadium	20-Apr-11	1107985-06	MW-3-110420
200.7	mg/L L	<0.00529	C	Thallium	20-Apr-11	1107985-06	MW-3-110420
200.7	mg/L L	81.0	D	Sodium	20-Apr-11	1107985-06	MW-3-110420
200.7	mg/L L	<0.00039	C,N	Silver	20-Apr-11	1107985-06	MW-3-110420
200.7	mg/L L	0.0337		Selenium	20-Apr-11	1107985-06	MW-3-110420
200.7	mg/L L	5.92	z	Potassium	20-Apr-11	1107985-06	MW-3-110420
200.7	mg/L L	<0.00116	C	Nickel	20-Apr-11	1107985-06	MW-3-110420
200.7	mg/L L	<0.00043	C	Manganese	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	57.5		Magnesium	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	<0.00210	C	Lead	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	<0.0106	C	Iron	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	<0.00183	C	Copper	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	<0.00051	C	Cobalt	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	<0.00109	C	Chromium	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	192	D	Calcium	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	<0.00082	C	Cadmium	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	<0.00072	C	Beryllium	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	0.0281	٦	Barium	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	<0.00206	C	Arsenic	20-Apr-11	1107985-06	MW-3-110420
	mg/L L	0.0024	ت	Antimony	20-Apr-11	1107985-06	MW-3-110420
<u>Method</u> 200.7	Units Matrix mg/L L	Result 0.0152	Flag J	<u>Analyte</u> Aluminum	Collected 20-Apr-11	Lab ID# 1107985-06	<u>Client ID#</u> MW-3-110420
					:		3::-:: 5:

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MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	<u>Client ID#</u> MW-4-110420
1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	1107985-07	<u>Lab ID#</u> 1107985-07
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	Collected 20-Apr-11
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
د	C	د	0	ب			J		D	ر	C	C	د	C	U	_	ر	ر	ر		Flag J
0.0103	<0.00174	0.0075	152	0.0025	0.0395	11.8	0.0022	0.0324	121	0.0027	<0.0106	<0.00183	0.0011	<0.00109	203	<0.00082	0.0008	0.0357	0.0041	0.0207	Result 0.0117
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	<u>Units</u> mg/L
· -		_	_	_	_	_	_	_	_	-	_	_	_	_	_	_	_	Г	_	г	Matrix L
200.7 1 0.00053 0.0200 29-Apr-11 VVK	200.7 1 0.00174 0.0200 29-Apr-11 VVK	200.7 1 0.00529 0.0200 29-Apr-11 VVK	200.7 1 0.197 2.00 11-May-11 VVK	200.7 1 0.00039 0.0200 29-Apr-11 VVK	200.7 1 0.00449 0.0200 29-Apr-11 VVK	200.7 1 0.0240 0.500 29-Apr-11 VVK	200.7 1 0.00116 0.0200 29-Apr-11 VVK	200.7 1 0.00043 0.0200 29-Apr-11 VVK	200.7 1 0.00079 0.100 11-May-11 VVK	200.7 1 0.00210 0.0400 29-Apr-11 VVK	200.7 1 0.0106 0.100 29-Apr-11 VVK	200.7 1 0.00183 0.0200 29-Apr-11 VVK	200.7 1 0.00051 0.0200 29-Apr-11 VVK	200.7 1 0.00109 0.0200 29-Apr-11 VVK	200.7 1 0.00393 0.500 11-May-11 VVK	200.7 1 0.00082 0.00500 29-Apr-11 VVK	200.7 1 0.00072 0.00400 29-Apr-11 VVK	200.7 1 0.00057 0.0400 29-Apr-11 VVK	200.7 1 0.00206 0.0200 29-Apr-11 VVK	200.7 1 0.00237 0.0200 29-Apr-11 VVK	Method DF LOD LOQ Run Analyst 200.7 1 0.0105 0.100 29-Apr-11 VVK

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MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	Client ID#						
1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	1107985-08	#CI 46.1
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	Callastad						
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	Analyte Aluminum	A 1
ے	_	ے	O	C			٦	<u>ر</u>	O	C	ے	C	_	C	D	C	C	د ـ	C	د	J	
0.0037	<0.00174	0.0081	450	0.0010	0.0288	16.9	0.0020	0.0193	205	<0.00210	0.0924	<0.00183	<0.00051	<0.00109	278	<0.00082	<0.00072	0.0148	<0.00206	0.0056	0.0161	Dh
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	1 1						
_	-	_	Г	Г	_	_	_	_	_	_	_	_	_		_	_	Г	Г	_	_	L	Matric
200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	Mathad
1 0.00053	1 0.00174	1 0.00529	1 0.197	1 0.00039	1 0.00449	1 0.0240	1 0.00116	1 0.00043	1 0.00079	1 0.00210	1 0.0106	1 0.00183	1 0.00051	1 0.00109	1 0.00393	1 0.00082	1 0.00072	1 0.00057	1 0.00206	1 0.00237	1 0.0105	
0.0200	0.0200	0.0200	2.00	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500	0.00500	0.00400	0.0400	0.0200	0.0200	0.100	-
29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 WK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 WK	11-May-11 VVK	29-Apr-11 WK	29-Apr-11 VVK									

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	53 0.0000	0.00053	200.7 1		ma/	0.0020	_	Zinc	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	74 0.0200	1 0.00174	200.7 1	F	mg/L	<0.00174		Vanadium	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	29 0.0200	1 0.00529	200.7 1	L 2	mg/L	0.0119	ے	Thallium	19-Apr-11	1107985-09	MW-7-110419
11-May-11 VVK	7 2.00	1 0.197	200.7 1	L 2	mg/L	289	D	Sodium	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	39 0.0200	1 0.00039	200.7 1	L	mg/L	<0.00039	_	Silver	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	49 0.0200	1 0.00449	200.7 1		mg/L	0.0296		Selenium	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	0.500	1 0.0240	200.7 1		mg/L	1.82		Potassium	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	16 0.0200	1 0.00116	200.7 1	_	mg/L	<0.00116	C	Nickel	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	43 0.0200	1 0.00043	200.7		mg/L	<0.00043	_	Manganese	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	79 0.100	1 0.00079	200.7		mg/L	1.64		Magnesium	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	10 0.0400	1 0.00210	200.7		mg/L	0.0029	ر	Lead	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK)6 0.100	1 0.0106	200.7	_	mg/L	0.0174	د	Iron	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	83 0.0200	1 0.00183	200.7		mg/L	<0.00183	C	Copper	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	51 0.0200	1 0.00051	200.7	F	mg/L	0.0006	ر	Cobalt	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	09 0.0200	1 0.00109	200.7	_	mg/L	<0.00109	C	Chromium	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	93 0.500	1 0.00393	200.7	_	mg/L	6.02	D	Calcium	19-Apr-11	1107985-09	MW-7-110419
0 29-Apr-11 VVK	0.00082 0.00500	1 0.000	200.7		mg/L	<0.00082	C	Cadmium	19-Apr-11	1107985-09	MW-7-110419
0 29-Apr-11 VVK	0.00072 0.00400	1 0.000	200.7	_	mg/L	<0.00072	C	Beryllium	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	57 0.0400	1 0.00057	200.7	_	mg/L	0.0274	ر	Barium	19-Apr-11	1107985-09	MW-7-110419
29-Apr-11 VVK	06 0.0200	1 0.00206	200.7	_	mg/L	0.0028	<u>د</u>	Arsenic	19-Apr-11	1107985-09	MW-7-110419
) 29-Apr-11 VVK	37 0.0200	1 0.00237	200.7	<u></u>	mg/L	0.0030	ے	Antimony	19-Apr-11	1107985-09	MW-7-110419
Run Analyst 29-Apr-11 VVK)5 <u>LOQ</u>	<u>DF</u> <u>LOD</u> 1 0.0105	Method 200.7	Matrix M	<u>Units</u> mg/L	Result 0.0286	Flag J	<u>Analyte</u> Aluminum	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-09	Client ID# MW-7-110419

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ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

29-Apr-11 VVK	0.0200	0.00053	р	200.7	۲	mg/L	<0.00053	C	Zinc	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00174	⊢	200.7	_	mg/L	<0.00174	_	Vanadium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00529	 	200.7	_	mg/L	<0.00529	_	Thallium	19-Apr-11	1107985-10	MW-8-110419
11-May-11 VVK	2.00	0.197	_	200.7	Г	mg/L	593	0	Sodium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00039	<u> </u>	200.7	_	mg/L	0.0011	Č, Z	Silver	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00449		200.7	_	mg/L	0.0317		Selenium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.500	0.0240	!t	200.7	Г	mg/L	5.59	z	Potassium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00116	_	200.7	_	mg/L	0.0019	ر	Nickel	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00043	\vdash	200.7	_	mg/L	<0.00043	C	Manganese	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.100	0.00079	<u></u>	200.7	_	mg/L	1.59		Magnesium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0400	0.00210	}	200.7	_	mg/L	<0.00210	C	Lead	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.100	0.0106	_	200.7	г	mg/L	0.0239	ت	Iron	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00183	<u></u>	200.7	_	mg/L	<0.00183	C	Copper	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00051	<u> </u>	200.7	_	mg/L	0.0010	ل	Cobalt	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00109		200.7	_	mg/L	0.0012	ے	Chromium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.500	0.00393	μ.	200.7	L	mg/L	15.3	D	Calcium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.00500	0.00082	<u> </u>	200.7	_	mg/L	<0.00082	C	Cadmium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.00400	0.00072	_	200.7	_	mg/L	<0.00072	C	Beryllium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0400	0.00057	_	200.7	_	mg/L	0.0389	ے	Barium	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00206	⊢ ⊸	200.7	_	mg/L	0.0033	د	Arsenic	19-Apr-11	1107985-10	MW-8-110419
29-Apr-11 VVK	0.0200	0.00237	_	200.7	_	mg/L	0.0096	ر	Antimony	19-Apr-11	1107985-10	MW-8-110419
Run Analyst 29-Apr-11 VVK	0.100	<u>LOD</u> 0.0105	DF 1	Method 200.7	Matrix L	<u>Units</u> mg/L	Result 0.0227	<u>Flag</u> J	<u>Analyte</u> Aluminum	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-10	<u>Client ID#</u> MW-8-110419

Phone: 330-253-8211

ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

MW-MD-02-110420	<u>Client ID#</u> MW-MD-02-110420																				
1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	1107985-11	<u>Lab ID#</u> 1107985-11
20-Apr-11	Collected 20-Apr-11																				
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
	C	ر	D	C			ر		D	C		C	٦	ے	D	C	C	د	ے	د	<u>Flag</u> U
<0.00053	<0.00174	0.0099	1410	<0.00039	0.0395	13.3	0.0057	0.693	252	<0.00210	1.61	<0.00183	0.0017	0.0014	214	<0.00082	<0.00072	0.0132	0.0102	0.0074	Result < 0.0105
mg/L	<u>Units</u> mg/L																				
	Г	Г	_	Г	_	-	<u></u>	_	_	_	٢	Г	_	_	Г	_	_	-	-	Г	Matrix L
200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	200.7 1	Method I 200.7 1
0.00053	0.00174	0.00529	0.197	0.00039	0.00449	0.0240	0.00116	0.00043	0.00079	0.00210	0.0106	0.00183	0.00051	0.00109	0.00393	0.00082	0.00072	0.00057 0.0400	0.00206	0.00237	<u>DF LOD</u> 1 0.0105
0.0200	0.0200	0.0200	2.00 1	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500	0.00500	0.00400	0.0400	0.0200	0.0200	<u>LOQ</u>
29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 WK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 WK	29-Apr-11 WK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	Run Analyst 29-Apr-11 VVK				

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

Received: 4/26/2011 Project #: Jackpile Dallas, TX 75354

	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MAN-MD-T10450	MW-MD-110420	Client ID#
				1107985-12	1107985-12	1107985-12				1107985-12					1107985-12	1107985-12	1107985-12	1107985-12	1107985-12	1107985-12	1107985-12	1107985-12	Lab ID#
- I	20-Apr-11	30-Apr 11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	Collected
	Zinc	Vandina	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	Aluminum	Analyte
	= c	: c	= -	O	C			د		0	C		C	ر	C	D	C	C	ر	ے	د	C	<u>Flag</u>
<0.0003	<0.00174	<0.00529		1310	<0.00039	0.0271	12.6	0.0048	0.654	237	<0.00210	1.45	<0.00183	0.0013	<0.00109	200	<0.00082	<0.00072	0.0116	0.0064	0.0041	<0.0105	Result
mg/L	mg/L	mg/L	: 'i'g/' r	ma/l	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	Units												
	<u> </u>	_	- r-	_	_	- -	_	_	_	_	_	-	_	_	_	_	_	r	_	_	_	_	Matrix
200.7 1	200.7 1	200.7 1	200./	7007	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	Method
0.00053 0.0200	0.00174	0.00529	. 0.197	0 107	0.00039	0.00449	0.0240	0.00116	1 0.00043	1 0.00079	1 0.00210	1 0.0106	1 0.00183	1 0.00051	1 0.00109	1 0.00393	1 0.00082	1 0.00073	1 0.00057	1 0.00206	1 0.00237	0	DF LOD
		0.0200	2.00	J (0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	3 0.500	0.00082 0.00500	0.00072 0.00400	0.00057 0.0400	5 0.0200	7 0.0200		LOQ
29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	11 N 11 N N N N N N N N N N N N N N N N	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	Run Analyst								

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

Lab Diff Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 1107985-13 20-Apr-11 Antimony U <0.00237 mg/L L 200.7 1 1107985-13 20-Apr-11 Berium U <0.00206 mg/L L 200.7 1 1107985-13 20-Apr-11 Berium J 0.0112 mg/L L 200.7 1 1107985-13 20-Apr-11 Cadmium U <0.00082 mg/L L 200.7 1 1107985-13 20-Apr-11 Cadmium D 571 mg/L L 200.7 1 1107985-13 20-Apr-11 Cabrit J 0.0604 mg/L L 200.7 1 1107985-13 20-Apr-11 Cabrit J 0.0692 mg/L L 200.7 1 1107985-	0.00053 0.0200 29-Apr-11 VVK	⊢ ;	200.7	F 1	mg/L	0.0107	: C	Zinc	20-Apr-11		NPOP20E-110420
Lab ID# Collected Analyze Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 1107985-13 20-Apr-11 Arsenic U <0.00237		<u> </u>	200.7		mg/L	<0.00529	C C	Vanadium	20-Apr-11	1107985-13	NPOP20E-110420
Lab LD# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum 1 0.8222 mg/L L 200.7 1 1107985-13 20-Apr-11 Arsenic U 0.00237 mg/L L 200.7 1 1107985-13 20-Apr-11 Arsenic U 0.00206 mg/L L 200.7 1 1107985-13 20-Apr-11 Bervillium U 0.0067 mg/L L 200.7 1 1107985-13 20-Apr-11 Cadmium U 0.0067 mg/L L 200.7 1 1107985-13 20-Apr-11 Cadmium U 0.00692 mg/L L 200.7 1 1107985-13 20-Apr-11 Cobalit J 0.0692 mg/L L 200.7 1 1107985-13 20-Apr-11 Cobalit J 0.0519			200.7		mg/L	1260	= 0	Sodium	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 1107985-13 20-Apr-11 Arsenic U <0.00237	0.00039 0.0200		200.7	_	mg/L	0.0522		Silver	20-Apr-11	1107985-13	NPOP20E-110420
Lab DB# Collected Analytic Flag Result Ubits Matrix Method DF 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 20.07 1 1107985-13 20-Apr-11 Antimony U <0.00237	0.00449 0.0200	<u> </u>	200.7	_	mg/L	0.0309		Selenium	20-Apr-11	1107985-13	NPOP20E-110420
Lab IDIIII Collected Analytic Flag Result Units Matin Method DF 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 1107985-13 20-Apr-11 Artimony U <0.00237	0.0240 0.500	<u> </u>	200.7	_	mg/L	39.7		Potassium	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L L L 200.7 1 1107985-13 20-Apr-11 Antimony U <0.00237	0.00116 0.0200	 	200.7	_	mg/L	0.0374		Nickel	20-Apr-11	1107985-13	NPOP20E-110420
Lab D# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 1107985-13 20-Apr-11 Antimony U <0.00237	0.00043 0.0200	<u> </u>	200.7	_	mg/L	4.633		Manganese	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 1107985-13 20-Apr-11 Antimony U <0.00237	0.00079 0.100	⊢	200.7	_	mg/L	276	O	Magnesium	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 1107985-13 20-Apr-11 Artimony U <0.00237	0.00210 0.0400	 4	200.7	_	mg/L	0.0519		Lead	20-Apr-11		NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum U <0.0227	0.0106 0.100	<u> </u>	200.7	_	mg/L	3.18		Iron	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum U <0.822	0.00183 0.0200	_	200.7	_	mg/L	<0.00183	C.	Copper	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum U <0.00237	0.00051 0.0200	ш	200.7	_	mg/L	0.0692	ت	Cobalt	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum U <0.0223	0.00109 0.0200	Н	200.7	_	mg/L	0.0604		Chromium	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum U <0.0237	0.00393 0.500	ш	200.7	_	mg/L	571	D	Calcium	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF 1107985-13 20-Apr-11 Aluminum U <0.0227	0.00082 0.00500	<u>—</u>	200.7	_	mg/L	<0.00082	_	Cadmium	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF LOD 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 0.0105 1107985-13 20-Apr-11 Antimony U <0.00237	0.00072 0.00400	<u> </u>	200.7	_	mg/L	0.0067		Beryllium	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF LOD 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 0.0105 1107985-13 20-Apr-11 Arsenic U <0.00206	0.00057 0.0400	<u> </u>	200.7	_	mg/L	0.0112	ر	Barium	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF LOD 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 0.00237 1107985-13 20-Apr-11 Antimony U <0.00237	0.00206 0.0200	_	200.7	_	mg/L	<0.00206	C	Arsenic	20-Apr-11	1107985-13	NPOP20E-110420
Lab ID# Collected Analyte Flag Result Units Matrix Method DF LOD 1107985-13 20-Apr-11 Aluminum 0.822 mg/L L 200.7 1 0.0105	0.00237 0.0200	_	200.7	_	mg/L	<0.00237	C	Antimony	20-Apr-11	1107985-13	NPOP20E-110420
			<u>Method</u> 200.7	Matrix L	Units mg/L	Result 0.822	Flag	<u>Analyte</u> Aluminum	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-13	Client ID# NPOP20E-110420

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

12-May-11 VVK	3.95	0.126	⊢	6010B	S	mg/Kg	65.0		Zinc	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.0790	↦	6010B	S	mg/Kg	44.8		Vanadium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.253	щ	6010B	S	mg/Kg	<0.253	C	Thallium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	79.0	68.7	<u> </u>	6010B	S	mg/Kg	5610		Sodium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.158		6010B	S	mg/Kg	<0.158	C	Silver	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.158	<u> </u>	6010 B	S	mg/Kg	<0.158	C	Selenium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	79.0	11.4	ᆫ	6010B	S	mg/Kg	9600		Potassium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.111	⊢	6010B	S	mg/Kg	14.7		Nickel	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.0948	<u>junt</u>	6010B	S	mg/Kg	558		Manganese	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	79.0	0.822	₩	6010B	S	mg/Kg	12300	D	Magnesium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	7.90	0.711	1	6010B	S	mg/Kg	9.69	د	Lead	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	79.0	1.01	1	6010B	S	mg/Kg	27300	D	Iron	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.126	Ľ	6010 B	S	mg/Kg	18.9		Copper	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.0948	\vdash	6010B	S	mg/Kg	7.30		Cobalt	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.0790	Ľ	6010 B	S	mg/Kg	17.7		Chromium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	79.0	1.07	1	6010B	S	mg/Kg	74200	D	Calcium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	0.790	0.0316	⊢	6010B	S	mg/Kg	0.901		Cadmium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.126	1	6010B	S	mg/ K g	1.19	<u>د</u>	Beryllium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	7.90	0.0474	H	6010B	S	mg/Kg	187		Barium	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3.95	0.300	Н	6010B	S	mg/Kg	7.82		Arsenic	20-Apr-11	1107985-14	PR-SED-01-110420
12-May-11 VVK	3,95	0.126	Н	6010B	S	mg/ K g	1.88	د	Antimony	20-Apr-11	1107985-14	PR-SED-01-110420
<u>Run</u> <u>Analyst</u> 12-May-11 VVK	<u>LOQ</u> 15.8	<u>LOD</u> 6.92	1 PF	<u>Method</u> 6010B	Matrix S	<u>Units</u> mg/Kg	<u>Result</u> 31900	<u>Flag</u> D	<u>Analyte</u> Aluminum	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-14	Client ID# PR-SED-01-110420

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ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

Client ID#

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

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Project #: Jackpile

29-Apr-11 VVK	0.0200	0.00053 0.0200	<u> </u>	200.7	L	mg/L	<0.00053	C	Zinc	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.0200	0.00174 0.0200		200.7	٢	mg/L	<0.00174	C	Vanadium	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.0200	0.00529 0.0200	-	200.7	_	mg/L	0.0130	۲	Thallium	20-Apr-11	1107985-15	110420
11-May-11 VVK	2.00	0.197		200.7	_	mg/L	1200	D	Sodium	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.0200	0.00039 0.0200	-	200.7	_	mg/L	<0.00039	C	Silver	20-Apr-11	1107985-15	110420
29-Apr-11 WK	0.0200	0.00449 0.0200	 	200.7	Г	mg/L	0.0302		Selenium	20-Apr-11	1107985-15	110420
11-May-11 VVK	0.500	0.0240	<u> </u>	200.7	_	mg/L	92.8	D	Potassium	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.0200	0.00116 0.0200	<u> </u>	200.7	-	mg/L	0.0032	د	Nickel	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.0200	0.00043	<u>,</u>	200.7	_	mg/L	0.970		Manganese	20-Apr-11	1107985-15	110420
11-May-11 VVK	0.100	0.00079	1	200.7	_	mg/L	471	D	Magnesium	20-Apr-11	1107985-15	110420
29-Apr-11 WK	0.0400	0.00210 0.0400	<u> </u>	200.7	_	mg/L	<0.00210	C	Lead	20-Apr-11	1107985-15	110420
29-Apr-11 WK	0.100	0.0106	<u> </u>	200.7	_	mg/L	0.269		Iron	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.0200	0.00183 0.0200	 	200.7	٢	mg/L	<0.00183	C	Copper	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.0200	0.00051 0.0200	<u> </u>	200.7	_	mg/L	0.0009	ر	Cobalt	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.0200	0.00109 0.0200	<u> </u>	200.7	_	mg/L	<0.00109	_	Chromium	20-Apr-11	1107985-15	110420
11-May-11 VVK	0.500	0.00393 0.500	_	200.7	_	mg/L	1010	D	Calcium	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.00500	0.00082 0.00500	₩	200.7	_	mg/L	<0.00082	C	Cadmium	20-Apr-11	1107985-15	110420
29-Apr-11 VVK	0.00400	0.00072 0.00400		200.7	_	mg/L	<0.00072	C	Beryllium	20-Apr-11	1107985-15	110420
29-Apr-11 WK	0.0400	0.00057 0.0400	_	200.7	_	mg/L	0.140		Barium	20-Apr-11	1107985-15	110420
29-Apr-11 WK	0.0200	0.00206 0.0200	_	200.7	_	mg/L	0.0075	ر	Arsenic	20-Apr-11	1107985-15	110420
29-Apr-11 WK	0.0200	0.00237 0.0200		200.7	_	mg/L	0.0033	ن	Antimony	20-Apr-11	1107985-15	110420
Run Analyst 29-Apr-11 VVK	0.100	1 0.0105	1	Method 200.7	Matrix L	Units mg/L	<u>Result</u> < 0.0105	<u>Flag</u> ∪	<u>Analyte</u> Aluminum	Collected 20-Apr-11	1107985-15	110420
) !) :	-	

PR-SW-01-1

PR-SW-01-1 PR-SW-01-1

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PR-SW-01-1: PR-SW-01-1 PR-SW-01-1 PR-SW-01-1: PR-SW-01-1 PR-SW-01-1 PR-SW-01-1 PR-SW-01-1 PR-SW-01-1

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011

Project #: Jackpile

RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	Client ID# RB-01-110420
1107985-16 20	1107985-16 20	1107985-16 20	1107985-16 20	1107985-16 20	1107985-16 20	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	1107985-16 2	<u>Lab ID#</u> <u>C</u> 1107985-16 2
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	Collected 20-Apr-11
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
۲	C	C	O	ر		C	د	C	⊂	ر	C	C	نــ	C	D	C	C	<u>_</u>	٦	<u>_</u>	Flag
0.0058	<0.00174	<0.00529	4.84	0.0029	0.0343	<0.0240	0.0012	<0.00043	<0.00079	0.0027	<0.0106	<0.00183	0.0006	<0.00109	3.60	<0.00082	<0.00072	0.0009	0.0033	0.0175	<u>Result</u> < 0.0105
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	Units mg/L
-	_	_	_	_	_	_	Г	_	Г-	_	_	_	_	_	_	٢	٢	Г	<u></u>	Г	Matrix L
200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	<u>Method</u> 200.7
1 0	1 0	1 0	₩.	1 C	1 0	_	C	1 C	1 C	1 C	_	1 (1 (1 (1 (1 (1 (1 (1 (1 (<u>DF</u>
0.00053 0.0200	0.00174	0.00529	0.197	0.00039	0.00449	0.0240	0.00116	0.00043	0.00079	0.00210	0.0106	0.00183	0.00051	0.00109	0.00393	0.00082	0.00072 (0.00057	0.00206	0.00237	<u>LOD</u> 0.0105
0.0200	0.0200	0.0200	2.00	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500	0.00500	0.00400	0.0400	0.0200	0.0200	<u>LOQ</u>
29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 WK	29-Apr-11 WK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 WK	29-Apr-11 WK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 WK	29-Apr-11 VVK	Run Analyst 29-Apr-11 VVK						

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June 06, 2011

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Received: 4/26/2011 Project #: Jackpile

RB-02-110421 1107985-17 21-Apr-11																					
. Zinc	l Vanadium	t Thallium	1 Sodium	1 Silver	1 Selenium	1 Potassium	1 Nickel	1 Manganese	1 Magnesium	1 Lead	1 Iron	1 Copper	1 Cobalt	1 Chromium	1 Calcium	1 Cadmium	1 Beryllium	1 Barium	1 Arsenic	1 Antimony	
<u>.</u>	_	۲	D	ر		\subset	C	\subset	O	C	C	ر	ر	د							
0.0067	<0.00174	0.0151	2.01	0.0012	0.0257	<0.0240	<0.00116	<0.00043	<0.00079	<0.00210	<0.0106	<0.00183	<0.00051	<0.00109	1.34	<0.00082	<0.00072	0.0011	0.0029	0.0076	
mg/L																					
	Γ	_	_	_	_	_	_	_	_	_	_	_	_	_	_	Γ	_	_	Г	_	
200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	
-	<u> </u>	<u> </u>	 	<u> </u>	_	المسا	Ľ	<u> </u>	 	1	_	Н	ᆫ	<u>ш</u>	↦	ᆢ	<u>ш</u>	 -	<u> </u>	<u> </u>	
0.00053 0.0200	0.00174	0.00529	0.197	0.00039	0.00449	0.0240	0.00116	0.00043	0.00079	0.00210	0.0106	0.00183	0.00051	0.00109	0.00393	0.00082 0.00500	0.00072	0.00057	0.00206	0.00237	
0.0200	0.0200	0.0200	2.00	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500	0.00500	0.00400	0.0400	0.0200	0.0200	
	29-Apr-11 WK	29-Apr-11 VVK																			

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| RM-JM-SED-02-110420 1107985-18 |
|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|
| | | | | | | | | | | | | | | | | | | | | | |
| 20-Apr-11 |
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	Aluminum
		\subset		C	C					ے	D				O	ت	ے			ر	O
34.2	19.8	<0.174	3880	<0.109	< 0.109	2310	9.36	160	5060	3.06	17900	8.27	5.22	7.88	20900	0.240	0.393	68.0	4.65	0.529	7420
mg/Kg	mg/Kg	mg/Kg	mg/ K g	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/ K g	mg/ K g	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/ K g	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/Kg
S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S
6010B	6010 B	6010B	6010B	6010B	6010B	6010 B	6010B	6010B	6010B	6010B											
<u>н</u>			<u> </u>	<u> </u>	Ľ	H	↦	<u> </u>	<u> </u>	1	_	Н		1	1	<u> </u>	⊢	H	<u> </u>	Н	щ
0.0872	0.0545	0.174	47.4	0.109	0.109	7.89	0.0763	0.0654	0.567	0.491	0.698	0.0872	0.0654	0.0545	0.741	0.0218	0.0872	0.0327	0.207	0.0872	4.78
2								N.i	ĹΠ	ū	ū	2	2.	2	Ų	0.	2	(n	ь т		
2 2.73	2.73	2.73	54.5	2.73	2.73	54.5	2.73	2.73	54.5	5.45	54.5	2.73	2.73	2.73	54.5	0.545	2.73	5.45	2.73	2.73	10.9

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RM-JM-SED-110420	Client ID# RM-JM-SED-110420																				
1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	1107985-19	<u>Lab ID#</u> 1107985-19
20-Apr-11	Collected 20-Apr-11																				
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
		ے	0	∪,N,*		z	z			د	0				D	C	ر				<u>Flag</u> D
36.0	22.4	0.717	4240	<0.119	23.3	2660	9.18	173	4890	2.35	16900	7.31	5.26	9.10	18000	<0.0237	0.558	57.0	5.72	23.3	Result 6660
mg/Kg	mg/ K g	mg/Kg	mg/ K g	mg/Kg	mg/Kg	mg/Kg	mg/ K g	mg/Kg	mg/Kg	mg/Kg	mg/Kg	<u>Units</u> mg/Kg									
S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	Matrix S
6010B	Method 6010B																				
1 0.0949	1 0.0593	0.190	1 51.6	0.119	0.119	1 8.59	1 0.0830	1 0.0712	1 0.617	1 0.534	1 0.759	1 0.0949	1 0.0712	1 0.0593	1 0.807	1 0.0237	1 0.0949	1 0.0356	1 0.225	1 0.0949	<u>DF LOD</u> 1 5.20
2.97	2.97	2.97	59.3	2.97	2.97	59.3	2.97	2.97	59.3	5.93	59.3	2.97	2.97	2.97	59.3	0.593	2.97	5.93	2.97	2.97	<u>LOQ</u> 11.9
29-Apr-11 VVK	29-Apr-11 WK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 WK	29-Apr-11 WK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	Run Analyst 11-May-11 VVK												

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Collected Analytic Flag Result Units Matrix Method DF LOD 20-Apr-11 Aluminum U 0.0105 mg/L L 200.0 1 0.0105 20.0 1 0.0105 20.0 1 0.0105 20.0 1 0.0105 20.0 1 0.0105 20.0 1 0.0105 20.00 1 0.0105 20.00 1 0.0105 20.00 1 0.00237 20.00 1 0.00237 20.00 1 0.00266 20.00 1 0.00267 1 0.00267 20.00 20.00 1 0.00057 1 0.00057 1 0.00057 20.00 20.00 1 0.00057 1 0.00057 1 0.00057 1 0.00057 1 0.00057 1 0.00058 20.00 1 0.00059 1 0.00057 1 0.00059 1 0.00059 1 0.00059 1 0.00059 0.00059 1 0.	29-Apr-11 VVK	0.0200	0.00053	1 (200.7	Г	mg/L	<0.00053		Zinc	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DF LOQ LOQ 20-Apr-11 Autminum U <0.0105	29-Apr-11 VVK	0.0200		1 (200.7	Г	mg/L	<0.00174	C	Vanadium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DF LOQ LOQ 20-Apr-11 Auminum U <0.0105	29-Apr-11 VVK	0.0200		1 (200.7	Г	mg/L	0.0154	ر	Thallium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method Dr. LOQ 20-Apr-11 Aluminum U <0.0105	11-May-11 VVK	2.00	0.197	<u> </u>	200.7	_	mg/L	532	D	Sodium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analytic Flag Result Units Matrix Method DF LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.0200		1 (200.7	_	mg/L	<0.00039	C	Silver	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analytic Flag Result Units Matrix Method DF LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.0200		1 (200.7	_	mg/L	0.0278		Selenium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Method DE LOD LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.500		<u> </u>	200.7	_	mg/L	18.6		Potassium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected 20-Apr-11 Analyte Aluminum Flag U Result CO.0105 Units mg/L Matrix L Method 2007 1 LOD 0.0105 LOD 0.1005 LOD 0.0105 LOD 0.0105 LOD 0.0105 Units 0.0105 Matrix 0.0105 Method 1 LOD 0.0105 LOD 0.0207 LOD 0.0207 <th< td=""><td>29-Apr-11 VVK</td><td>0.0200</td><td></td><td>1 (</td><td>200.7</td><td>_</td><td>mg/L</td><td>0.0023</td><td>ر</td><td>Nickel</td><td>20-Apr-11</td><td>1107985-20</td><td>RM-JM-SW-02</td></th<>	29-Apr-11 VVK	0.0200		1 (200.7	_	mg/L	0.0023	ر	Nickel	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analytic Flag Result Units Matrix Method DE LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.0200		1 (200.7	_	mg/L	0.0152	C	Manganese	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analytic Flag Result Units Matrix Method DF LOQ 20-Apr-11 Aluminum U <0.0105	11-May-11 VVK	0.100	ე.00079	1 (200.7	_	mg/L	222	0	Magnesium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DE LOQ LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.0400		1 (200.7	_	mg/L	<0.00210	_	Lead	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DE LOD LOD 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.100		H	200.7	_	mg/L	<0.0106	C	Iron	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DE LOD LOD 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.0200		1 (200.7	г	mg/L	<0.00183	\subset	Copper	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DE LOD LOD 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.0200		1 (200.7	_	mg/L	<0.00051	_	Cobalt	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DE LOD LOD 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.0200		1 (200.7	_	mg/L	<0.00109	C	Chromium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DE LOD LOD 20-Apr-11 Aluminum U <0.0105	11-May-11 VVK	0.500			200.7	_	mg/L	224	D	Calcium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DE LOD LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.00500	0.00082	1 (200.7	_	mg/L	<0.00082	_	Cadmium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DF LOD LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 WK	0.00400		1 (200.7		mg/L	<0.00072	C	Beryllium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DF LOD LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.0400		1 (200.7	٦	mg/L	0.0232	ے	Barium	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DF LOD LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 WK	0.0200		1	200.7	Г	mg/L	<0.00206	C	Arsenic	20-Apr-11	1107985-20	RM-JM-SW-02
Collected Analyte Flag Result Units Matrix Method DF LOD LOQ 20-Apr-11 Aluminum U <0.0105	29-Apr-11 VVK	0.0200		1	200.7	_	mg/L	0.0062	ے	Antimony	20-Apr-11	1107985-20	RM-JM-SW-02
	Run Analyst 29-Apr-11 VVK	<u>LOQ</u>	<u>LOD</u> 0.0105		Method 200.7	Matrix L	Units mg/L	Result < 0.0105	Flag U	<u>Analyte</u> Aluminum	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-20	Client ID# RM-JM-SW-02

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

RM-JM-SW-110420	Client ID# RM-JM-SW-110420																				
1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	1107985-21	<u>Lab ID#</u> 1107985-21
20-Apr-11	Collected 20-Apr-11																				
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
C	C	د	D	C			_	ر	D	ے	C	C	C	_	D	C	C	٦	_	د	Flag U
<0.00053	<0.00174	0.0175	519	<0.00039	0.0324	18.1	<0.00116	0.0131	217	0.0025	<0.0106	<0.00183	<0.00051	<0.00109	218	<0.00082	<0.00072	0.0228	<0.00206	0.0041	<u>Result</u> < 0.0105
mg/L	<u>Units</u> mg/L																				
	_	_	_	_	Г	_	Г	_	_	Г	_	_	_	_	_	_	_	-	Г	_	Matrix L
200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	Method 200.7
1 0	1 0	1 0	1	1 0	1 0	1 (1 0	1 0	1 0	1 0	1 -	1 0	1 0	1 C	1 C	1 0	1 (1 (1 (1 (DF 1
0.00053	0.00174 (0.00529 (0.197	0.00039	0.00449 (0.0240	0.00116 (0.00043 (0.00079	0.00210 (0.0106	0.00183 (0.00051 (0.00109 (0.00393	0.00082 0.00500	0.00072 0	0.00057	0.00206 (0.00237	<u>LOD</u> 0.0105
0.0200	0.0200	0.0200	2.00	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500).00500	0.00400	0.0400	0.0200	0.0200	<u>LOQ</u> 0.100
29-Apr-11 WK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	Run Analyst 29-Apr-11 VVK												

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29-Apr-11 VVK	3.52	0.113	⊢	6010B	S	mg/Kg	57.0		Zinc	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.0704	ь.	6010B	S	mg/Kg	29.8		Vanadium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.225	 	6010B	S	mg/Kg	0.662	د	Thallium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	70.4	61.3	<u>⊷</u>	6010B	S	mg/Kg	1950	O	Sodium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.141	<u> </u>	6010B	S	mg/ K g	< 0.141	C	Silver	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.141	<u></u>	6010 B	S	mg/Kg	29.1		Selenium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	70.4	10.2		6010B	S	mg/Kg	4180		Potassium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.0986		6010B	S	mg/Kg	14.6		Nickel	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.0845	H	6010B	S	mg/Kg	226		Manganese	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	70.4	0.732	<u> </u>	6010B	S	mg/Kg	6910		Magnesium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	7.04	0.634	_	6010B	S	mg/Kg	4.58	د	Lead	19-Apr-11	1107985-22	RM-SED-BG-110419
11-May-11 VVK	70.4	0.902		6010 B	S	mg/Kg	25900	O	Iron	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.113	 4	6010B	S	mg/Kg	13.1		Copper	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.0845	<u> </u>	6010B	S	mg/ K g	7.78		Cobalt	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.0704	_	6010B		mg/Kg	8.40		Chromium	19-Apr-11	1107985-22	RM-SED-BG-110419
11-May-11 VVK	70.4	0.958	<u> </u>	6010B	S	mg/Kg	23300	0	Calcium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	0.704	0.0282	Н	6010B	S	mg/Kg	<0.0282	C	Cadmium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.113	⊢	6010B	S	mg/Kg	0.916	د	Beryllium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	7.04	0.0423	1	6010B	S	mg/Kg	154		Barium	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.268	Н	6010B	S	mg/Kg	6.68		Arsenic	19-Apr-11	1107985-22	RM-SED-BG-110419
29-Apr-11 VVK	3.52	0.113	Ľ	6010B	S	mg/ K g	< 0.113	C	Antimony	19-Apr-11	1107985-22	RM-SED-BG-110419
Run Analyst 11-May-11 VVK	<u>LOQ</u> 14.1	<u>LOD</u> 6.17	<u>DF</u> 1	Method 6010B	Matrix S	<u>Units</u> mg/Kg	<u>Result</u> 15800	Flag D	<u>Analyte</u> Aluminum	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-22	Client ID# RM-SED-BG-110419

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

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29-Apr-11 VVK	0.0200	0.00053 0.0200	-	200.7	- -	mg/L	0.0023		Zinc	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00174	-	200.7	٢	mg/L	<0.00174	C	Vanadium	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00529	 	200.7	_	mg/L	0.0151	۲.	Thallium	19-Apr-11	1107985-23	RM-SW-BG-110419
11-May-11 VVK	2.00	0.197	<u> </u>	200.7	_	mg/L	280	D	Sodium	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00039	<u> </u>	200.7	_	mg/L	<0.00039	C	Silver	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00449	<u> </u>	200.7	_	mg/L	0.0350		Selenium	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.500	0.0240	<u> </u>	200.7	-	mg/L	15.0		Potassium	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00116	_	200.7	_	mg/L	<0.00116	C	Nickel	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00043	⊢	200.7	_	mg/L	<0.00043	C	Manganese	19-Apr-11	1107985-23	RM-SW-BG-110419
11-May-11 VVK	0.100	0.00079	<u> </u>	200.7	_	mg/L	106	0	Magnesium	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0400	0.00210	<u> </u>	200.7	_	mg/L	<0.00210	C	Lead	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.100	0.0106	р., .	200.7	_	mg/L	<0.0106	C	Iron	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00183	<u> </u>	200.7	_	mg/L	<0.00183	C	Copper	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00051	<u> </u>	200.7	_	mg/L	<0.00051	C	Cobalt	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00109		200.7	_	mg/L	<0.00109	C	Chromium	19-Apr-11	1107985-23	RM-SW-BG-110419
11-May-11 VVK	0.500	0.00393		200.7	_	mg/L	117	Ū	Calcium	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.00500	0.00082 0.00500	_	200.7	_	mg/L	<0.00082	C	Cadmium	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.00400	0.00072 0.00400	_	200.7	_	mg/L	<0.00072	C	Beryllium	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0400	0.00057 0.0400	⊢	200.7	٦	mg/L	0.0261	ر	Barium	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00206	<u>_</u>	200.7	_	mg/L	<0.00206	C	Arsenic	19-Apr-11	1107985-23	RM-SW-BG-110419
29-Apr-11 VVK	0.0200	0.00237	_	200.7	_	mg/L	0.0043	ے	Antimony	19-Apr-11	1107985-23	RM-SW-BG-110419
Run Analyst 29-Apr-11 VVK	0.100	0.0105	<u>DF</u>	<u>Method</u> 200.7	Matrix L	Units mg/L	<u>Result</u> <0.0105	Flag U	<u>Analyte</u> Aluminum	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-23	Client ID# RM-SW-BG-110419

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01	RP-JM-SED-01							
1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24	1107985-24
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11							
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	Aluminum
		_	D	\subset						ے	D				D	C	_			C	ם
34.1	23.8	<0.187	2330	<0.117	30.0	1950	10.6	490	4650	1.90	32600	6.18	6.29	9.33	30400	<0.0234	0.615	116	6.57	<0.0937	7230
mg/Kg	mg/ K g	mg/Kg	mg/Kg	mg/ K g	mg/Kg	mg/ K g	mg/Kg	mg/ K g	mg/Kg	mg/Kg	mg/ K g	mg/Kg	mg/Kg	mg/Kg	mg/Kg						
S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S
6010B	6010B	6010B	6010 B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B
1 0	1 0	_	щ	_	_	⊢	1 (1 (<u> </u>	<u> </u>	<u> </u>	1 (1 (1 (<u> </u>	1 (1 (1 (1	<u></u>	_
0.0937	0.0585	0.187				8.48	0.0819	0.0702					0.0702	0.0585	0.796	,			0.222	0.0937	5.13
2.93	2.93	2.93	58.5	2.93	2.93	58.5	2.93	2.93	58.5	5.85	58.5	2.93	2.93	2.93	58.5	0.585	2.93	5.85	2.93	2.93	11.7
29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VV	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK							

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

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	RP-JM-SED-110420	RP-JM-SFD-110420	RP-JM-SED-110420	Client ID# RP-JM-SED-110420																		
			1107985-25	1107985-25	1107985-25									1107985-25			1107985-25		1107985-25	1107985-25	1107985-25	<u>Lab ID#</u> 1107985-25
1 0 1 1	20-Apr-11	20-Apr-11	20-Anr-11	20-Apr-11	Collected 20-Apr-11																	
	Zinc	Vandium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
	C	: c	Ξ	Ū	C						د	D				D	C	ر			C	<u>Flag</u> D
32.3	24.3	21.2 <0.210	100	1520	<0.131	23.6	3410	11.1	166	6430	3.15	19800	9.81	5.63	11.7	18200	<0.0263	0.729	72.9	5.73	<0.105	<u>Result</u> 13500
mg/kg	mg/Kg	mg/kg		ma/ K a	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/ K g	mg/Kg	<u>Units</u> mg/Kg										
v	, v	· v) (S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	S	Matrix S
6010B	6010B	9010B		6010B	Method 6010B																	
–	1 (٠ ٢	-1	} ≜	—	Ľ	_	_	1		⊢	_		_	<u> </u>	<u>⊢</u>	jk	 	 	-	1 <u>D</u> F
0.105	•			57 1	0.131	0.131	9.51	0.0919	-	0.683	0.591	0.841	0.105	0.0788	0.0657	0.893	0.0263	0.105	0.0394	0.250	0.105	<u>LOD</u> 5.75
3.28		3.28	00.7	4 ع ا	3.28	3.28	65.7			65.7	6.57	65.7	3.28	3.28	3.28	65.7	0.657	3.28	6.57	3.28	3.28	<u>LOQ</u> 13.1
29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	23-ADI-11 VVN	20-Apr 11 10/V	29-Apr-11 VVK	29-Apr-11 WK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 WK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	Run <u>Analyst</u> 11-May-11 VVK					

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RP-JM-SW-01-110420	RP-JM-SW-01-1104:20	RP-JM-SW-01-1104:20	RP-JM-SW-01-110420	<u>Client ID#</u> RP-JM-SW-01-110420																	
1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	1107985-26	<u>Lab ID#</u> 1107985-26
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	Collected 20-Apr-11							
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
· · ·	C	٦	D	C			C		D	C	C	C	C	C	D	C	C	د	C	C	Flag U
0.0025	<0.00174	0.0071	295	<0.00039	0.0341	15.3	<0.00116	0.0348	150	<0.00210	<0.0106	<0.00183	<0.00051	<0.00109	200	<0.00082	<0.00072	0.0370	<0.00206	<0.00237	Result < 0.0105
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	<u>Units</u> mg/L							
	Г	_	Г			_	_	_	Γ	_	_	_	_	_	_	_	٦	Г	Г	Г	Matrix L
200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	Method 200.7
1 0.0	1 0.0	1 0.0	1 0	1 0.0	1 0.0	1 0.	1 0.0	1 0.0	1 0.0	1 0.0	1 0.	1 0.0	1 0.0	1 0.0	1 0.0	1 0.0	1 0.0	1 0.0	1 0.	1 0.	<u>DF</u> .
0.00053 0.0200	0.00174 0.0200	0.00529	0.197	0.00039	0.00449	0.0240	0.00116	0.00043	0.00079	0.00210	0.0106	0.00183	0.00051	0.00109	0.00393	00082	00072	00057	0.00206	0.00237	<u>LOD</u> 0.0105
0.0200	0.0200	0.0200	2.00	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500	0.00082 0.00500	0.00072 0.00400	0.00057 0.0400	0.0200	0.0200	<u>LOQ</u> 0.100
29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 WK	29-Apr-11 WK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 WK	29-Apr-11 WK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	Run Analyst 29-Apr-11 VVK

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

RP-JM-SW-110420	Client ID# RP-JM-SW-110420																				
1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	1107985-27	<u>Lab ID#</u> 1107985-27
20-Apr-11	Collected 20-Apr-11																				
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
ے	C	C	D	د			ر		D	C	C	C	C	C	Ū	_	C		C	ر	<u>Flag</u>
0.0034	<0.00174	<0.00529	127	0.0012	0.0276	12.3	0.0019	0.0374	96.7	<0.00210	<0.0106	<0.00183	<0.00051	<0.00109	181	<0.00082	<0.00072	0.0472	<0.00206	0.0048	<u>Result</u> < 0.0105
mg/L	<u>Units</u> mg/L																				
	_	_	_	_	_	_	_	_	_	_	_	Г	٦	_	٦	_	Г	Γ-	_	Г	Matrix L
200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	Method 200.7
1 0	1 0	1 (1 (1 0	_	1 (1	1 (1 (—	1 (1 (<u> </u>	1 (1 (1 (1 (<u> </u>	1 (<u>DF</u> 1
0.00053	0.00174 0.0200	0.00529	0.197	0.00039	0.00449	0.0240	0.00116	0.00043	0.00079	0.00210	0.0106	0.00183	0.00051	0.00109	0.00393	0.00082 0.00500	0.00072 0.00400	0.00057 0.0400	0.00206	0.00237	<u>LOD</u> 0.0105
0.0200	0.0200	0.0200	2.00	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500	0.00500	0.00400	0.0400	0.0200	0.0200	<u>LOQ</u>
29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	Run Analyst 29-Apr-11 VVK												

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June 06, 2011

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Received: 4/26/2011 Project #: Jackpile

29-Apr-11 VVK 29-Apr-11 VVK	3.19	0.102										
29-Apr-11 VVK		0 102	_	6010B	S	mg/Kg	32.7		Zinc	19-Apr-11	1107985-28	RP-SED-01-110419
29-ADI-11 VVN	3.19	0.0638	⊢	6010B	S	mg/Kg	18.6		Vanadium	19-Apr-11	1107985-28	RP-SED-01-110419
JO 12 11 WW	3.19	0.204	-	6010B	S	mg/Kg	0.747	٦	Thallium	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	63.8	55.5	<u> </u>	6010B	S	mg/Kg	4830	D	Sodium	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	3.19	0.128	<u> </u>	6010B	S	mg/Kg	<0.128	U	Silver	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	3.19	0.128	 	6010B	S	mg/Kg	18.4		Selenium	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	63.8	9.24	<u> </u>	6010B	S	mg/Kg	4060		Potassium	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	3.19	0.0893	1	6010B	S	mg/Kg	7.54		Nickel	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	3.19	0.0766	⊢	6010B	S	mg/Kg	150		Manganese	19-Apr-11	1107985-28	RP-SED-01-110419
11-May-11 VVK	63.8	0.664	 	6010B	S	mg/Kg	8410	D	Magnesium	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	6.38	0.574	Ľ	6010B	S	mg/Kg	3.32	٦	Lead	19-Apr-11	1107985-28	RP-SED-01-110419
11-May-11 VVK	63.8	0.817	Н	6010 B	S	mg/Kg	15500	D	Iron	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 WK	3.19	0.102	⊷	6010B	S	mg/Kg	7.26		Copper	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	3.19	0.0766	Ľ	6010B	S	mg/ K g	4.00		Cobalt	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	3.19	0.0638	—	6010B	S	mg/Kg	8.09		Chromium	19-Apr-11	1107985-28	RP-SED-01-110419
11-May-11 VVK	63.8	0.868	<u>ш</u>	6010B	S	mg/Kg	26000	D	Calcium	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 WK	0.638	0.0255	<u> </u>	6010B	S	mg/ K g	<0.0255	C	Cadmium	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 WK	3.19	0.102	H	6010B	S	mg/Kg	0.574	Ĵ	Beryllium	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	6.38	0.0383	<u> </u>	6010B	S	mg/Kg	76.2		Barium	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 WK	3.19	0.243	<u> </u>	6010B	S	mg/Kg	4.56		Arsenic	19-Apr-11	1107985-28	RP-SED-01-110419
29-Apr-11 VVK	3.19	0.102	-	6010B	S	mg/Kg	<0.102	C	Antimony	19-Apr-11	1107985-28	RP-SED-01-110419
Run Analyst 11-May-11 VVK	<u>LOQ</u> 12.8	<u>LOD</u> 5.59	<u>DF</u>	<u>Method</u> 6010B	Matrix S	<u>Units</u> mg/Kg	<u>Result</u> 10100	<u>Flag</u> D	<u>Analyte</u> Aluminum	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-28	<u>Client ID#</u> RP-SED-01-110419

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

29-Apr-11 VVK	3.22	0.103	\vdash	6010B	S	mg/Kg	38.8		Zinc	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 WK	3.22	0.0644	_	6010B	S	mg/Kg	22.2		Vanadium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.206	<u> </u>	6010B	S	mg/Kg	<0.206	C	Thallium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	64.4	56.0	↦	6010B	S	mg/Kg	1850	D	Sodium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.129	<u></u>	6010B	S	mg/Kg	<0.129	_	Silver	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.129	щ	6010B	S	mg/Kg	19.9		Selenium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	64.4	9.33	<u> </u>	6010B	S	mg/Kg	4250		Potassium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.0902	ㅂ	6010B	S	mg/Kg	8.95		Nickel	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.0773	<u> </u>	6010B	S	mg/Kg	258		Manganese	20-Apr-11	1107985-29	RP-SED-02-110420
11-May-11 VVK	64.4	0.670		6010B	S	mg/Kg	7920	D	Magnesium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	6.44	0.580	н	6010B	S	mg/Kg	2.59	J	Lead	20-Apr-11	1107985-29	RP-SED-02-110420
11-May-11 VVK	64.4	0.825	<u> </u>	6010B	S	mg/Kg	16900	D	Iron	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.103	⊢	6010B	S	mg/Kg	8.87		Copper	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.0773	<u>-</u>	6010B	S	mg/Kg	4.88		Cobalt	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.0644	ш	6010 B	S	mg/Kg	9.58		Chromium	20-Apr-11	1107985-29	RP-SED-02-110420
11-May-11 VVK	64.4	0.876	μ	6010B	S	mg/ K g	31700	D	Calcium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	0.644	0.0258	Н	6010B	S	mg/Kg	<0.0258	C	Cadmium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.103	⊢	6010B	S	mg/Kg	0.728	ے	Beryllium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	6.44	0.0386	<u> </u>	6010B	S	mg/ K g	157		Barium	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.245	ш	6010B	S	mg/Kg	4.52		Arsenic	20-Apr-11	1107985-29	RP-SED-02-110420
29-Apr-11 VVK	3.22	0.103	<u></u>	6010B	S	mg/Kg	<0.103	C	Antimony	20-Apr-11	1107985-29	RP-SED-02-110420
Run Analyst 11-May-11 VVK	<u>LOQ</u> 12.9	<u>LOD</u> 5.64	<u>DF</u>	<u>Method</u> 6010B	Matrix S	<u>Units</u> mg/Kg	Result 13900	<u>Flag</u> D	<u>Analyte</u> Aluminum	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-29	Client ID# RP-SED-02-110420

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Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

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| RP-SED-03-110420 1107985-30 |
|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|
| 20-Apr-11 | 20-Apr-11 | 20-Apr-11 | 20-Apr-11 | 20-Apr-11 |) 20-Apr-11 | 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |) 20-Apr-11 |
| Zinc | Vanadium | Thallium | Sodium | Silver | Selenium | Potassium | Nickel | Manganese | Magnesium | Lead | Iron | Copper | Cobalt | Chromium | Calcium | Cadmium | Beryllium | Barium | Arsenic | Antimony |
| | | C | O | _ | | | | | O | ر | 0 | | | | D | C | ے | | | C |
| 60.1 | 34.6 | <0.168 | 1450 | <0.105 | 28.7 | 5460 | 14.7 | 231 | 9800 | 3.22 | 25200 | 14.1 | 7.17 | 16.1 | 26900 | <0.0209 | 1.14 | 132 | 6.05 | <0.0838 |
| mg/ K g | mg/ K g | mg/Kg | mg/Kg | mg/Kg | mg/Kg | mg/Kg | mg/ K g | mg/Kg |
| S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S | S |
| 6010B |
⊢	1	_	—	<u> </u>	_	⊢ •	<u> </u>	—	—	<u> </u>	⊢	<u> </u>	<u> </u>	 	-	<u></u>	-	<u> </u>	<u></u>	_
0	0	0.	45.6	0.105	0.105	7.58	0.0733	0.0628	0.545	0.471	0.670	0.0838	0.0628	0.0524	0.712	0.0209	0.0838	0.0314	0.199	0.0838
 0.0838	0.0524	0.168	.6	5	ഗ്	•	ω	œ												
.0838 2.62	0524 2.62	168 2.62	.6 52.4)5 2.62)5 2.62	52.4	3 2.62	8 2.62	52.4	5.24	52.4	2.62	2.62	2.62	52.4	0.524	2.62	5.24	2.62	2.62

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

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RP-SED-BG-041911 1107985-31 19-Apr-11	1107985-31	RP-SED-BG-041911 1107985-31 19-Apr-11	NF-3ED-BG-041911 110/963-31 19-Api-11		1107985-31	1107985-31 1107985-31	1107985-31 1107985-31 1107985-31	1107985-31 1107985-31 1107985-31 1107985-31	1107985-31 1107985-31 11107985-31 11107985-31 11107985-31	1107985-31 1107985-31 1107985-31 1107985-31 1107985-31 1107985-31	1107985-31 1107985-31 1107985-31 1107985-31 1107985-31 1107985-31	1107985-31 1107985-31 1107985-31 1107985-31 1107985-31 1107985-31 1107985-31	1107985-31 1107985-31 1107985-31 1107985-31 1107985-31 1107985-31 1107985-31 1107985-31								
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead		Iron	Copper Iron	Cobalt Copper Iron	Chromium Cobalt Copper Iron	Calcium Chromium Cobalt Copper					
	ı	\subset	D	C						د		O	O	O	Ū	0 0	ם סכ	0 0 0 0	0 0 0 0	ם סכי	0 0 0 0 0 0
38.7	29.9	<0.213	1040	< 0.133	28.3	2170	9.17	358	3880	1.72	2000	32800	5.94 32800	9.34 5.94 32800	9.12 9.34 5.94	28500 9.12 9.34 5.94	<0.0266 28500 9.12 9.34 5.94	0.586 <0.0266 28500 9.12 9.34 5.94	97.7 0.586 <0.0266 28500 9.12 9.34 5.94	4.53 97.7 0.586 <0.0266 28500 9.12 9.34 5.94	<0.106 4.53 97.7 0.586 <0.0266 28500 9.12 9.34 5.94
mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/ K g	mg/Kg	mg/Kg	mg/Kg	mg/ K g	mg/Kg	ì	mg/Kg	mg/Kg mg/Kg	mg/Kg mg/Kg mg/Kg	mg/Kg mg/Kg mg/Kg mg/Kg	mg/Kg mg/Kg mg/Kg mg/Kg mg/Kg	mg/Kg mg/Kg mg/Kg mg/Kg mg/Kg	mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg	mg/Kg mg/Kg mg/Kg mg/Kg mg/Kg mg/Kg mg/Kg	mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg
	S	S	S	S	S	S	S	S	S	S	S		S	S S	у у у	у у у у	w w w w	w w w w w			w w w w w w w
			C				•	C	Ф.	_	0		6								
S 6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B	6010B		010B	5010B 5010B	5010B 5010B 5010B	6010B 6010B 6010B	6010B 6010B 6010B 6010B	6010B 6010B 6010B 6010B 5010B	5010B 5010B 5010B 5010B 5010B 5010B	5010B 5010B 5010B 5010B 5010B 5010B 5010B	6010B 6010B 6010B 6010B 6010B 6010B
H	⊢ +	<u></u>		⊢	ь		1	⊢	H	₽	—		 								
1 0.106	1 0.0665	1 0.213	1 57.9	1 0.133	1 0.133	1 9.64	1 0.0932	1 0.0799	1 0.692	1 0.599	1 0.852		1 0.106	1 0.0799 1 0.106	1 0.0665 1 0.0799 1 0.106	1 0.905 1 0.0665 1 0.0799 1 0.106	1 0.0266 1 0.905 1 0.0665 1 0.0799 1 0.106	1 0.106 1 0.0266 1 0.905 1 0.0665 1 0.0799 1 0.106	1 0.0399 1 0.106 1 0.0266 1 0.905 1 0.0665 1 0.0799	1 0.253 1 0.0399 1 0.106 1 0.0266 1 0.905 1 0.0665 1 0.0799	1 0.106 1 0.253 1 0.0399 1 0.106 1 0.0266 1 0.905 1 0.0665 1 0.0799
1 0.106	1 0.0665	1 0.213		⊢	ь		1	⊢	H	₽	—		1 0.106	1 0.0799 1 0.106	1 0.0665 1 0.0799 1 0.106	1 0.905 1 0.0665 1 0.0799 1 0.106	1 0.0266 1 0.905 1 0.0665 1 0.0799 1 0.106	1 0.106 1 0.0266 1 0.905 1 0.0665 1 0.0799 1 0.106	1 0.0399 1 0.106 1 0.0266 1 0.905 1 0.0665 1 0.0799	1 0.253 1 0.0399 1 0.106 1 0.0266 1 0.905 1 0.0665 1 0.0799	1 0.106 1 0.253 1 0.0399 1 0.106 1 0.0266 1 0.905 1 0.0665 1 0.0799

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

29-Apr-11 VVK	0.0200	0.00053	 	200.7	_	mg/L	<0.00053	C	Zinc	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00174	1	200.7	_	mg/L	<0.00174	C	Vanadium	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00529	μ	200.7	_	mg/L	0.0082	۲	Thallium	19-Apr-11	1107985-32	RP-SW-01-110419
11-May-11 VVK	2.00	0.197	<u> </u>	200.7	_	mg/L	445	D	Sodium	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00039	 4	200.7	_	mg/L	<0.00039	C	Silver	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00449	<u> </u>	200.7	_	mg/L	0.0299		Selenium	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.500	0.0240	<u> </u>	200.7	_	mg/L	17.3		Potassium	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00116	1-4	200.7	_	mg/L	0.0025	د ــ	Nickel	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00043	<u> </u>	200.7	_	mg/L	0.0424		Manganese	19-Apr-11	1107985-32	RP-SW-01-110419
11-May-11 VVK	0.100	0.00079	<u>-</u>	200.7	_	mg/L	211	D	Magnesium	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0400	0.00210		200.7	_	mg/L	<0.00210	C	Lead	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.100	0.0106	<u> </u>	200.7	_	mg/L	<0.0106	□	Iron	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00183	<u> </u>	200.7	_	mg/L	<0.00183	C	Copper	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00051	⊢	200.7	_	mg/L	0.0007	د	Cobalt	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00109	<u> </u>	200.7	г-	mg/L	<0.00109	U	Chromium	19-Apr-11	1107985-32	RP-SW-01-110419
11-May-11 VVK	0.500	0.00393	⊢	200.7	_	mg/L	223	D	Calcium	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.00500	0.00082		200.7	Г	mg/L	<0.00082	C	Cadmium	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.00400	0.00072	<u> </u>	200.7	_	mg/L	<0.00072	C	Beryllium	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0400	0.00057	<u></u>	200.7	_	mg/L	0.0255	٦	Barium	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00206	⊨	200.7	٦	mg/L	<0.00206	C	Arsenic	19-Apr-11	1107985-32	RP-SW-01-110419
29-Apr-11 VVK	0.0200	0.00237	⊢	200.7	L	mg/L	0.0047	ے	Antimony	19-Apr-11	1107985-32	RP-SW-01-110419
Run Analyst 29-Apr-11 VVK	<u>LOQ</u>	<u>LOD</u> 0.0105	<u>DF</u>	<u>Method</u> 200.7	Matrix L	Units mg/L	Result < 0.0105	<u>Flag</u> U	<u>Analyte</u> Aluminum	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-32	Client ID# RP-SW-01-110419

ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911
1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33	1107985-33
19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	Aluminum
	_		_	_																	ı
J	C	ت	D	\subset			\subset			\subset	\subset	\subset	\subset		D	\subset	\subset			<u> </u>	-
0.0046	<0.00174	0.0111	57.3	<0.00039	0.0273	6.16	<0.00116	0.641	36.0	<0.00210	<0.0106	<0.00183	<0.00051	<0.00109	124	<0.00082	<0.00072	0.0742	<0.00206	<0.00237	<0.0105
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	3	_					_	_	_	_	_			
		Г	٦	_		_	/	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L
_	۳		ر د	L	_	_	/L	g/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L	ng/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L	mg/L L
L 200.7	L 200.7	L L 200.7	'L L 200.7	L L 200.7	L L 200.7	L L 200.7	/L L 200.7	g/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	ng/L L 200.7	ng/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	mg/L L 200.7	
ь	<u>-</u>	L 200.7 1	_	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	Г	L 200.7 1	_	-	٢	_	_	L 200.7 1	L 200.7 1	—	-
1 0.00053	1 0.00174	L 200.7 1 0.00529	_	L 200.7 1 0.00039	L 200.7 1 0.00449	L 200.7 1 0.0240	L 200.7 1 0.00116	L 200.7 1 0.00043	L 200.7 1 0.00079	L 200.7 1 0.00210	L 200.7 1 0.0106	L 200.7 1 0.00183	L 200.7 1 0.00051	L 200.7 1 0.00109	L 200.7 1 0.00393	L 200.7 1 0.00082	L 200.7 1	L 200.7 1	L 200.7 1 0.00206	L 200.7 1 0.00237	L 200.7
ь	<u>-</u>	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	L 200.7 1	_	г	L 200.7 1	L 200.7 1	L 200.7 1

Phone: 330-253-8211

ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	<u>Client ID#</u> SPOP35-110421
1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	1107985-34	<u>Lab ID#</u> 1107985-34
21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	Collected 21-Apr-11
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
_	C	C	O	ن			ر		D	ت	ے	C	د	٦	D	C	C	ے	C	C	<u>Flag</u>
<0.00053	<0.00174	<0.00529	2030	0.0047	0.0337	63.5	0.0062	0.186	359	0.0080	0.0504	<0.00183	0.0016	0.0057	483	<0.00082	<0.00072	0.0070	<0.00206	<0.00237	Result 0.0237
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	<u>Units</u> mg/L
_	r 	-	_	<u>-</u>	Г	Г	_	_	Г	_	_	_	_	_	_	Г	_	_	Г	_	Matrix L
200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	Method 200.7
	<u> </u>	<u> </u>	⊢	<u> </u>	-	 1	⊢	⊢	 4	⊢	⊣	⊢ ~	<u> </u>	<u> </u>	ب	<u> </u>	<u> </u>	<u> </u>	 	_	<u>DF</u>
0.00053 0.0200	0.00174	0.00529	0.197	0.00039	0.00449	0.0240	0.00116	0.00043	0.00079	0.00210	0.0106	0.00183	0.00051	0.00109	0.00393	0.00082 0.00500	0.00072	0.00057	0.00206	0.00237	<u>LOD</u> 0.0105
	0.0200	0.0200	2.00	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500	0.00500	0.00400	0.0400	0.0200	0.0200	<u>LOQ</u> 0.100
29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 WK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	11-May-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 VVK	29-Apr-11 WK	Run Analyst 29-Apr-11 VVK				

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ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

MW-RM-110420	MW-RM-110420	MW-RM-110420	MW-RM-110420	<u>Client ID#</u> MW-RM-110420																	
1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	1107985-35	<u>Lab ID#</u> 1107985-35
21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	Collected 21-Apr-11																	
Zinc	Vanadium	Thallium	Sodium	Silver	Selenium	Potassium	Nickel	Manganese	Magnesium	Lead	Iron	Copper	Cobalt	Chromium	Calcium	Cadmium	Beryllium	Barium	Arsenic	Antimony	<u>Analyte</u> Aluminum
C	C	٦	D	_			C	C	D	C	C	<u>_</u>	C	C	D	_	C	د	C	<u>_</u>	Flag
<0.00053	<0.00174	0.0082	187	<0.00039	0.0344	19.4	<0.00116	<0.00043	111	<0.00210	<0.0106	0.0029	<0.00051	<0.00109	146	<0.00082	<0.00072	0.0134	<0.00206	0.0114	<u>Result</u> 0.278
mg/L	mg/L	mg/L	mg/L	Units mg/L																	
. -	_	_	_	г	г	_	_	Г	_	_	_	_	_	_	_	_	г	_	_	٢	Matrix L
200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	200.7	<u>Method</u> 200.7
1 (1 (1 (μ.		1 (<u> </u>	1	⊢		1	<u> </u>		1		_	-	_	<u>-</u>	<u>-</u>		
0.00053	0.00174	0.00529	0.197	0.00039	0.00449	0.0240	0.00116	0.00043	0.00079	0.00210	0.0106	0.00183	0.00051	0.00109	0.00393	0.00082	0.00072 0.00400	0.00057	0.00206	0.00237	<u>LOD</u> 0.0105
0.0200	0.0200	0.0200	2.00	0.0200	0.0200	0.500	0.0200	0.0200	0.100	0.0400	0.100	0.0200	0.0200	0.0200	0.500	0.00500	0.00400	0.0400	0.0200	0.0200	<u>LOQ</u>
12-May-11 VVK	12-May-11 VVK	12-May-11 VVK	12-May-11 VVK	Run Analyst 12-May-11 VVK																	

Phone: 330-253-8211

ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

<u>DF LOD LOQ</u> 1 0.74 1 DF LOD LOQ
DF DF

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ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

MW-1-110419	Client ID#
1107985-03	Lab ID#
19-Apr-11	Collected
1107985-03 19-Apr-11 Gross Alpha	Analyte
<u> </u>	Result
13.4 +/-1.5	sult
pci/l	Units
٦	Matrix
SM 7110C	Method
<u></u>	<u>P</u>
2.28	LOD
ω	<u>100</u>
05-May-11 MO	Run Analyst

~	→	
<u>Client ID#</u> MW-1-110419	9	<u>Client ID#</u> MW-1-110419
<u>Lab ID#</u> 1107985-03	- 6-	<u>Lab ID#</u> <u>Collec</u> 1107985-03 19-Ap
Collected 19-Apr-11	Collected 19-Apr-11	<u>ted</u> r-11
Collected Analyte 3 19-Apr-11 Radium-228	<u>Analyte</u> Radium-226	Analyte Gross Beta
<u>Result</u> 1.26 +/-0.4	<u>Result</u> U +/-0.24	Result U +/-3.97
Units pci/l	Units pci/l	Units pci/l
Matrix L	Matrix L	Matrix L
Method DF LOD EPA904.0 1 0.92	Method EPA 903.0	Method <u>DF</u> <u>LOD</u> EPA 900.0 1 3.58
1	<u>DF</u> <u>LOD</u> 1 0.74	DF
<u>LOD</u> 0.92	<u>LOD</u> 0.74	<u>LOD</u> 3.58
1 100	<u>LOQ</u>	<u>LOQ</u>
Run Analyst 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analys 29-Apr-11 MO

1107985-04	Lab ID#	
21-Apr-11	Collected	
107985-04 21-Apr-11 Gross Alpha	<u>Analyte</u>	
65.5 +/-4.1	Result	
pci/I	Units	
_	Matrix	
SM 7110C	Method	
1	DF	
2.28	<u>LOD</u>	
 ω		
05-May-11 MO	Run Analyst	

<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-	1107985-	1107985-04
<u>Coll</u>	<u>Collected</u>	<u>Collected</u>
21-A	04 21-Apr-11	04 21-Apr-11
<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>
Radium-226	Gross Beta	Gross Alpha
	<u>.</u>	6.
<u>sult</u> J +/-0.39	<u>sult</u> 9.3 +/-8.6	Result 65.5 +/-4.1
Units	Units	Units
pci/I	pci/l	pci/l
Matrix	Matrix	Matrix
L	L	L
Method	Method	<u>Method</u>
EPA 903.0	EPA 900.0	SM 7110C
<u>DF</u>	DF 1	<u>DF</u>
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
0.74	3.58	2.28
<u>LOQ</u>	100 4	3 100
<u>Run</u> <u>Anal</u>	Run <u>Anal</u>	Run Analyst
04-May-11 M	29-Apr-11 M	05-May-11 MO
	Collected Analyte Result Units Matrix Method DF LOD -04 21-Apr-11 Radium-226 U +/-0.39 pci/l L EPA 903.0 1 0.74	Collected Analyte Result Units Matrix Method DF LOD LOQ -04 21-Apr-11 Gross Beta 99.3 +/-8.6 pci/l L EPA 900.0 1 3.58 4 Collected Analyte Result Units Matrix Method DF LOD LOQ -04 21-Apr-11 Radium-226 U +/-0.39 pci/l L EPA 903.0 1 0.74 1

Client ID# MW-2-110421

MW-2-110421

Client ID#

MW-2-110421

1107985-04 21-Apr-11 Radium-228

U +/-0.36

pci/I

EPA904.0

<u>,</u>

0.92

<u>...</u>

03-May-11 MO

Client ID#

MW-2-110421

Client ID#

ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

Client ID#	<u>Client ID#</u>	<u>Client ID#</u>	Client ID#	Client ID#	Client ID#	Client ID#	Client ID#
MW-3-110420	MW-3-110420	MW-3-110420	MW-3-110420	MW-2-2-110421	MW-2-2-110421	MW-2-2-110421	MW-2-2-110421
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-06	1107985-06	1107985-06	1107985-06	1107985-05	1107985-05	1107985-05	1107985-05
Collected	Collected	Collected	Collected	Collected	Collected	Collected	Collected
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11
Analyte	Analyte	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	Analyte	<u>Analyte</u>
Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	Gross Alpha
Result	Result U +/-0.24	Result	Result	<u>Result</u>	<u>Result</u>	Result	Result
U +/-0.53		5.3 +/-1.7	43 +/-3.3	U +/-0.32	U +/-0.27	120.3 +/-9.4	70 +/-4.2
Units	Units	Units	Units	Units	Units	Units	Units
pci/I	pci/l	pci/l	pci/l	pci/l	pci/l	pci/l	pci/l
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L	L	L	L	L	L	L
Method	Method	Method	Method	Method	Method	Method	<u>Method</u>
EPA904.0	EPA 903.0	EPA 900.0	SM 7110C	EPA904.0	EPA 903.0	EPA 900.0	SM 7110C
<u>DF</u>	DF 1	1 DF	DF 1	<u>DF</u>	<u>DF</u>	DF	<u>DF</u>
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
0.92	0.74	3.58	2.28	0.92	0.74	3.58	2.28
<u> 1</u>	1	4	<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u>
Run Analyst 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 29-Apr-11 MO	Run Analyst 05-May-11 MO	Run <u>Analyst</u> 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 29-Apr-11 MO	Run Analyst 05-May-11 MO

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ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

Client ID#	<u>Client ID#</u>	Client ID#	<u>Client ID#</u>	Client ID#	<u>Client ID#</u>	Client ID#	<u>Client ID#</u>
MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-08	1107985-08	1107985-08	1107985-08	1107985-07	1107985-07	1107985-07	1107985-07
Collected	Collected	Collected	Collected	Collected	Collected	Collected	Collected
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
<u>Analyte</u>	Analyte	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>
Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	Gross Alpha
Result 2.51 +/-0.65	Result U +/-0.29	Result 66.7 +/-4.9	Result 61.8 +/-3.9	Result U +/-0.29	Result U +/-0.27	Result 91.3 +/-4.14	Result 142 +/-5.9
Units	Units	<u>Units</u>	<u>Units</u>	Units	Units	Units	Units
pci/I	pci/I	pci/I	pci/l	pci/l	pci/l	pci/l	pci/l
Matrix	<u>Matrix</u>	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L	L	L	L	L	L	L
Method	Method	Method	Method	Method	Method	Method	Method
EPA904.0	EPA 903.0	EPA 900.0	SM 7110C	EPA904.0	EPA 903.0	EPA 900.0	SM 7110C
DF DF	DF 1	<u>DF</u>	DF 1	DF	<u>DF</u>	DE 1	DF
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
0.92	0.74	3.58	2.28	0.92	0.74	3.58	2.28
1 00	<u>LOQ</u> 1	<u>LOQ</u>	<u>ΓΟΩ</u> 3	<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u> 4	<u>LOQ</u>
Run Analyst 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 29-Apr-11 MO	Run Analyst 05-May-11 MO	Run <u>Analyst</u> 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 29-Apr-11 MO	

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ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	Client ID#	Client ID#	<u>Client ID#</u>	<u>Client ID#</u>
MW-8-110419	MW-8-110419	MW-8-110419	MW-8-110419	MW-7-110419	MW-7-110419	MW-7-110419	MW-7-110419
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-10	1107985-10	1107985-10	1107985-10	1107985-09	1107985-09	1107985-09	1107985-09
Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11
<u>Analyte</u>	Analyte	Analyte	Analyte	Analyte	Analyte	Analyte	Analyte
Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	Gross Alpha
Result U +	Result U+/	<u>Result</u> 224 +	Result 40.5	Result 1.11 +	Result 2.71 +	Result U+	Result 1047 +
<u>esult</u>	<u>\(\csult\) +/-0.27</u>	Result	<u>Result</u>	Result	Result	<u>esult</u>	Result
U +/-0.3		224 +/-6.88	40.5 +/-3.4	1.11 +/-0.40	2.71 +/-0.55	U +/-1.2	1047 +/-15.9
<u>Units</u>	Units	Units	Units	Units	Units	Units	<u>Units</u>
pci/l	pci/l	pci/l	pci/1	pci/l	pci/l	pci/I	pci/l
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L	L	L	L	L	L	L
Method	Method	Method	Method	Method	Method	Method	<u>Method</u>
EPA904.0	EPA 903.0	EPA 900.0	SM 7110C	EPA904.0	EPA 903.0	EPA 900.0	SM 7110C
1 P	DF 1	DF 1	1 1	DF 1	DF 1	<u>DF</u>	<u>DF</u>
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
0.92	0.74	3.58	2.28	0.92	0.74	3.58	2.28
<u>L00</u>	<u> 1</u>	<u>LOQ</u>	3 3	<u>LOQ</u> 1	1	<u>LOQ</u>	<u>LOQ</u>
<u>Run Analyst</u> 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 29-Apr-11 MO	Run Analyst 05-May-11 MO	Run Analyst 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 29-Apr-11 MO	Run Analyst 05-May-11 MO

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Client ID#	Client ID#	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	Client ID#	<u>Client ID#</u>	Chent II)#
MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-02-110420	MW-MD-02-110420	MW-MD-02-110420	MW-MD-02-110420
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-12	1107985-12	1107985-12	1107985-12	1107985-11	1107985-11	1107985-11	1107985-11
Collected 20-Apr-11	Collected	Collected	Collected	Collected	Collected	Collected	Collected
	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
Analyte Radium-228	<u>Analyte</u> Radium-226	<u>Analyte</u> Gross Beta	Analyte Gross Alpha	<u>Analyte</u> Radium-228	<u>Analyte</u> Radium-226	<u>Analyte</u> Gross Beta	Analyte Gross Alpha
Result U +/-0.29	Result 1.34 +/-0.44	Result 17.8 +/-2.9	Result 27 +/-2.6	Result U +/-0.32	<u>Result</u> U +/-0.31	Result 20.7 +/-11.18	<u>Result</u> 149 +/-6
Units	Units	Units	<u>Units</u>	<u>Units</u>	<u>Units</u>	<u>Units</u>	Units
pci/I	pci/l	pci/l	pci/l	pci/I	pci/l	pci/l	pci/1
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L	L	L	L	L	L	L
Method	Method	Method	Method	Method	Method	Method	<u>Method</u>
EPA904.0	EPA 903.0	EPA 900.0	SM 7110C	EPA904.0	EPA 903.0	EPA 900.0	SM 7110C
<u>DF</u>	1 <u>DF</u>	<u>DF</u>	DF 1	DF	<u>DF</u>	1	1 1
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
0.92	0.74	3.58	2.28	0.92	0.74	3.58	2.28
1	1 1	<u>LOQ</u>	<u>LOQ</u> 3	<u> </u>	<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u> 3
Run Analyst 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 30-Apr-11 MO	Run Analyst 05-May-11 MO	Run Analyst 03-May-11 MO	<u>Run Analyst</u> 04-May-11 MO	Run Analyst 11-May-11 MO	Run Analyst 05-May-11 MO

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PR-SED-01-110420	PR-SED-01-110420	PR-SED-01-110420	PR-SED-01-110420	NPOP20E-110420	NPOP20E-110420	NPOP20E-110420	NPOP20E-110420
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-14	1107985-14	1107985-14	1107985-14	1107985-13	1107985-13	1107985-13	1107985-13
Collected	Collected	Collected	Collected	Collected	Collected	Collected	Collected
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
<u>Analyte</u>	Analyte	Analyte	Analyte	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	Analyte
Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	Gross Alpha
Result 3.40 +/-0.72	Result 3.17 +/-0.59	Result 9.6 +/-0.15	Result 4.83 +/-0.18	Result 13.93 +/-1.48	Result 33.23 +/-1.74	Result 26301.04 +/-277.55	<u>Result</u> 3624 +/-29
Units	Units	<u>Units</u>	Units	Units	<u>Units</u>	<u>Units</u>	Units
pci/g	pci/g	pci/g	pci/g	pci/I	pci/l	pci/l	pci/l
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
S	S	S	S	L	L	L	L
<u>Method</u>	<u>Method</u>	<u>Method</u>	<u>Method</u>	<u>Method</u>	Method	Method	Method
9320	9315	9310	9310	EPA904.0	EPA 903.0	EPA 900.0	SM 7110C
<u>DF</u>	<u>DF</u>	DF 1	DF	<u>DF</u>	<u>DF</u>	1 1	1 <u>P</u> F
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
2	1	3	2	0.92	0.74	3.58	2.28
<u> 100</u> 2	1 1	3	<u>LOQ</u>	1	<u>LOQ</u>	4	3
Run Analyst	Run Analyst	Run Analyst	Run Analyst	Run Analyst 03-May-11 MO	Run Analyst	Run <u>Analyst</u>	Run Analyst
04-May-11 MO	04-May-11 MO	05-May-11 MO	05-May-11 MO		04-May-11 MO	11-May-11 MO	05-May-11 MO

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<u>Client ID# Lab ID# Collected Analyte</u> RB-01-110420 1107985-16 20-Apr-11 Radium-228	Client ID# Lab ID# Collected Analyte RB-01-110420 1107985-16 20-Apr-11 Radium-226	Client ID# Lab ID# Collected Analyte RB-01-110420 1107985-16 20-Apr-11 Gross Beta	Client ID# Lab ID# Collected Analyte RB-01-110420 1107985-16 20-Apr-11 Gross Alpha	Client ID# <u>Lab ID#</u> <u>Collected</u> <u>Analyte</u> PR-SW-01-110420 1107985-15 20-Apr-11 Radium-228	<u>Client ID# Lab ID# Collected Analyte</u> PR-SW-01-110420 1107985-15 20-Apr-11 Radium-226	Client ID# Lab ID# Collected Analyte PR-SW-01-110420 1107985-15 20-Apr-11 Gross Beta	<u>Client ID#</u> <u>Lab ID#</u> <u>Collected</u> <u>Analyte</u> PR-SW-01-110420 1107985-15 20-Apr-11 Gross Alpha
Result Units 1.07 +/-0.4 pci/l	<u>Result</u> <u>Units</u> U +/-0.2 pci/l	<u>Result</u> <u>Units</u> 38.2 +/-2.67 pci/l	Result Units 26.7 +/-2.6 pci/l	Result <u>Units</u> U +/-0.27 pci/l	<u>Result</u> <u>Units</u> U +/-0.2 pci/l	Result Units 151.8 +/-5.6 pci/l	<u>Result</u> 55 +/-3.8 pci/l
its Matrix	its Matrix	iis Matrix	ils Matrix	its <u>Matrix</u> I L	<u>its Matrix</u>	its <u>Matrix</u>	<u>its</u> <u>Matrix</u> I L
Method <u>DF</u> <u>LOD</u> EPA904.0 1 0.92	Method <u>DF</u> <u>LOD</u> EPA 903.0 1 0.74	Method DF LOD EPA 900.0 1 3.58	Method DF LOD SM 7110C 1 2.28	Method <u>DF LOD</u> EPA904.0 1 0.92	Method DF LOD EPA 903.0 1 0.74	Method DF LOD EPA 900.0 1 3.58	Method DF LOD SM 7110C 1 2.28
<u>LOQ</u> 1	1	<u>LOQ</u>	3	1 1	<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u>
Run Analyst 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 30-Apr-11 MO	Run Analyst 05-May-11 MO	Run Analyst 03-May-11 MO	Run <u>Analyst</u> 04-May-11 MO	Run Analyst 30-Apr-11 MO	Run Analyst 05-May-11 MO

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Client ID# RM-JM-SED-02- 110420	Client ID# RM-JM-SED-02- 110420	Client ID# RM-JM-SED-02- 110420	Client ID# RB-02-110421	Client ID# RB-02-110421	Client ID# RB-02-110421	<u>Client ID#</u> RB-02-110421
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-18	1107985-18	1107985-18	1107985-17	1107985-17	1107985-17	1107985-17
Collected	Collected	Collected	Collected	Collected	Collected	Collected
20-Apr-11	20-Apr-11	20-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11
Analyte	Analyte	<u>Analyte</u>	Analyte	<u>Analyte</u>	<u>Analyte</u>	Analyte Gross Alpha
Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	
	: : : : : : : : : : :	<u>R.</u>	<u>R</u>			:
Result 1 +/-0.39	Result U +/-0.1	Result 2.43 +/-0.15	Result U +/-0.34	Result U +/-0.26	Result U +/-1.12	Result U +/-1.1
<u>Units</u>	<u>Units</u>	Units	Units	Units	Units	Units
pci/g	pci/g	pci/g	pci/l	pci/l	pci/I	pci/l
Matrix	Matrix	Matrix	Matrix	<u>Matrix</u>	Matrix	Matrix
S	S	S	L	L	L	L
<u>Method</u>	<u>Method</u>	<u>Method</u>	Method	Method	Method	<u>Method</u>
9315	9310	9310	EPA904.0	EPA 903.0	EPA 900.0	SM 7110C
<u>DF</u>	1 DF	<u>DF</u>	<u>DF</u>	<u>DF</u>	<u>DF</u>	DE
<u>LOD</u>	LOD	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
1	3		0.92	0.74	3.58	2.28
1 1	3	<u>LOQ</u> 2	<u>LOQ</u>	1 1	<u>LOQ</u> 4	<u>LOQ</u>
Run <u>Analyst</u> 04-May-11 MO	Run <u>Analyst</u> 05-May-11 MO	Run Analyst 05-May-11 MO	Run Analyst 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 30-Apr-11 MO	Run Analyst 05-May-11 MO

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Client ID# RM-JM-SW-02	Client ID# RM-JM-SW-02	Client ID# RM-JM-SW-02	Client ID# RM-JM-SED-110420	Client ID# RM-JM-SED-110420	Client ID# RM-JM-SED-110420	Client ID# RM-JM-SED-110420	<u>Client ID#</u> RM-JM-SED-02- 110420
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-20	1107985-20	1107985-20	1107985-19	1107985-19	1107985-19	1107985-19	1107985-18
Collected	Collected	Collected	Collected	Collected	Collected	Collected	Collected
20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
Analyte	Analyte	Analyte	Analyte	Analyte	Analyte	Analyte	<u>Analyte</u>
Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228
Result U +/-0.29	<u>Result</u> 39.4 +/-5.8	Result 23.6 +/-2.5	<u>Result</u> U +/-0.51	<u>Result</u> U +/-0.26	Result 9.75 +/-0.14	Result 5.45 +/-0.19	Result 2.64 +/-0.63
Units	Units	Units	Units	Units	Units	<u>Units</u>	Units
pci/l	8 pci/l	5 pci/I	pci/g	pci/g	L4 pci/g	19 pci/g	53 pci/g
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L	L	S	S	S	S	S
Method DF LOD	Method <u>DF</u> <u>LOD</u>	Method DF LOD	Method DF LOD 9320 1 2	Method DF LOD	Method <u>DF</u> <u>LOD</u>	Method <u>DF</u> <u>LOD</u>	Method <u>DF</u> <u>LOD</u>
EPA 903.0 1 0.74	EPA 900.0 1 3.58	SM 7110C 1 2.28		9315 1 1	9310 1 3	9310 1 2	9320 1 2
<u>LOQ</u>	<u>LOQ</u>	<u> </u>	<u>) LOQ</u> 2	<u>) LOQ</u> 1	<u>) LOQ</u>	<u> </u>	<u>) LOQ</u> 2
Run Analyst 04-May-11 MO	Run Analyst 30-Apr-11 MO	Run Analyst 05-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 05-May-11 MO	Run Analyst 05-May-11 MO	Run <u>Analyst</u> 04-May-11 MO

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<u>Client ID#</u>	Client ID#	Client ID#	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	Client ID#
RM-SED-BG-110419	RM-SED-BG-110419	RM-SED-BG-110419	RM-JM-SW-110420	RM-JM-SW-110420	RM-JM-SW-110420	RM-JM-SW-110420	RM-JM-SW-02
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-22	1107985-22	1107985-22	1107985-21	1107985-21	1107985-21	1107985-21	1107985-20
Collected 19-Apr-11	Collected	Collected	Collected	Collected	Collected	Collected	Collected
	19-Apr-11	19-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
<u>Analyte</u>	Analyte	<u>Analyte</u>	<u>Analyte</u>	Analyte	<u>Analyte</u>	Analyte	Analyte
Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228
Result 1.17 +/-0.4	Result 4.1 +/-0.11	Result 4.8 +/-0.19	Result U +/-0.4	Result U +/-0.32	<u>Result</u> 45.6 +/-5.65	Result 28.2 +/-2.8	Result 2.02 +/-0.6
<u>Units</u>	Units	Units	<u>Units</u>	Units	Units	Units	<u>Units</u>
pci/g	pci/g	pci/g	pci/l	pci/l	pci/I	pci/l	pci/l
Matrix	Matrix	Matrix	<u>Matrix</u>	Matrix	Matrix	Matrix	Matrix
S	S	S	L	L	L	L	L
Method	<u>Method</u>	<u>Method</u>	Method	Method	<u>Method</u>	Method	Method
9315	9310	9310	EPA904.0	EPA 903.0	EPA 900.0	SM 7110C	EPA904.0
DF	<u>DF</u>	<u>DF</u>	<u>DF</u>	1 <u>DF</u>	<u>DF</u>	1 DF	DF
LOD	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
1		2	0.92	0.74	3.58	2.28	0.92
<u>LOQ</u> 1	<u> </u>	<u>LOQ</u> 2	<u>Год</u> 1	<u>LOQ</u> 1	<u>LOQ</u>	<u>LOQ</u> 3	1 1
Run Analyst	Run Analyst	Run Analyst	Run Analyst 02-May-11 MO	Run Analyst	Run <u>Analyst</u>	Run <u>Analyst</u>	Run Analyst
04-May-11 MO	05-May-11 MO	05-May-11 MO		04-May-11 MO	30-Apr-11 MO	05-May-11 MO	02-May-11 MO

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Client ID# RP-JM-SED-01	Client ID# RP-JM-SED-01	Client ID# RP-JM-SED-01	Client ID# RM-SW-BG-110419	Client ID# RM-SW-BG-110419	Client ID# RM-SW-BG-110419	Client ID# RM-SW-BG-110419	Client ID# RM-SED-BG-110419
<u>Lab ID#</u> 1107985-24	<u>Lab ID#</u> 1107985-24	<u>Lab ID#</u> 1107985-24	<u>Lab ID#</u> 1107985-23	<u>Lab ID#</u> 1107985-23	<u>Lab ID#</u> 1107985-23	<u>Lab ID#</u> 1107985-23	<u>Lab ID#</u> 1107985-22
Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11
Analyte Radium-226	Analyte Gross Beta	Analyte Gross Alpha	<u>Analyte</u> Radium-228	<u>Analyte</u> Radium-226	Analyte Gross Beta	<u>Analyte</u> Gross Alpha	<u>Analyte</u> Radium-228
<u>Result</u> ∪ +/-0.37	Result 1.99 +/-0.1	Result 2.8 +/-0.15	Result 1.12 +/-0.45	Result U +/-0.34	Result 5.6 +/-2.46	Result U +/-1.2	Result U +/-0.48
Units pci/g	Units pci/g	Units pci/g	Units pci/l	Units pci/l	Units pci/1	<u>Units</u> pci/l	<u>Units</u> pci/g
Matrix S	Matrix S	Matrix S	Matrix L	<u>Matrix</u> L	Matrix L	Matrix L	Matrix S
<u>Method</u> 9315	Method 9310	<u>Method</u> 9310	Method EPA904.0	Method EPA 903.0	Method EPA 900.0	Method SM 7110C	<u>Method</u> 9320
DF	<u>DF</u>	<u>DF</u>	<u>DF</u>	DF 1	<u>DF</u>	<u>DF</u>	1 <u>DF</u>
<u>LOD</u> 1	3	<u>LOD</u> 2	<u>LOD</u> 0.92	<u>LOD</u> 0.74	<u>LOD</u> 3.58	<u>LOD</u> 2.28	<u>LOD</u> 2
<u> </u>	<u>100</u>	<u>LOQ</u>	<u>Loo</u>	1 LOQ	<u>LOQ</u>	<u>100</u>	<u>LOQ</u> 2
Run Analyst 04-May-11 MO	Run Analyst 05-May-11 MO	Run Analyst 05-May-11 MO	Run Analyst 02-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 30-Apr-11 MO	Run Analyst 05-May-11 MO	Run Analyst 04-May-11 MO

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

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<u>Client ID#</u> RP-JM-SW-01- 110420	<u>Client ID#</u> RP-JM-SW-01- 110420	Client ID# RP-JM-SED-110420	Client ID# RP-JM-SED-110420	Client ID# RP-JM-SED-110420	<u>Client ID#</u> RP-JM-SED-110420	Client ID# RP-JM-SED-01
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-26	1107985-26	1107985-25	1107985-25	1107985-25	1107985-25	1107985-24
Collected	Collected	Collected	Collected 20-Apr-11	Collected	Collected	Collected
20-Apr-11	20-Apr-11	20-Apr-11		20-Apr-11	20-Apr-11	20-Apr-11
Analyte	Analyte	Analyte	<u>Analyte</u>	Analyte	Analyte	<u>Analyte</u>
Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228
Result 25.9 +/-3.7	Result 30.8 +/-2.8	Result U +/-0.14	Result 1.77 +/-0.47	Result 2.4 +/-0.1	Result 4.6 +/-0.21	Result 2.14 +/-0.57
<u>Units</u>	<u>Units</u>	<u>Units</u>	Units pci/g	Units	Units	Units
pci/l	pci/l	pci/g		pci/g	pci/g	pci/g
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L	S	S	S	S	S
Method	Method	<u>Method</u>	<u>Method</u>	<u>Method</u>	Method	Method
EPA 900.0	SM 7110C	9320	9315	9310	9310	9320
1 DF	DF I	1 PF	DF I	1 DF	<u>DF</u> !	DF .
<u>LOD</u> 3.58	<u>LOD</u> 2.28	<u>LOD</u> 2	<u>LOD</u>	<u>LOD</u>	<u>LOD</u> 2	<u>LOD</u> 2
<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u> 2	1 1	3	<u>LOQ</u> 2	<u>LOQ</u> 2
Run <u>Analyst</u>	Run <u>Analyst</u>	Run Analyst	Run Analyst	Run Analyst	Run <u>Analyst</u> 05-May-11 MO	Run Analyst
30-Apr-11 MO	05-May-11 MO	04-May-11 MO	04-May-11 MO	05-May-11 MO		04-May-11 MO

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Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

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<u>Client ID#</u> RP-SED-01-110419	<u>Client ID#</u> RP-SED-01-110419	<u>Client ID#</u> RP-JM-SW-110420	Client ID# RP-JM-SW-110420	<u>Client ID#</u> RP-JM-SW-110420	Client ID# RP-JM-SW-110420	<u>Client ID#</u> RP-JM-SW-01- 110420	Client ID# RP-JM-SW-01- 110420
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-28	1107985-28	1107985-27	1107985-27	1107985-27	1107985-27	1107985-26	1107985-26
Collected 19-Apr-11	Collected 19-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11
<u>Λnalyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>
Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226
<u>Re</u>	$\frac{\mathrm{Re}}{1}$	<u> </u>	Re	<u>Re</u> 23	Re	Re Re	Re Re
Result 1.13 +/-0.13	Result 1.5 +/-0.15	Result U +/-0.35	Result U +/-0.32	Result 23.7 +/-2.77	Result 35.1 +/-3	Result U +/-0.38	Result U +/-0.35
Units	Units	Units	Units	Units	Units	<u>Units</u>	Units
pci/g	pci/g	pci/l	pci/l	pci/l	pci/I	pci/l	pci/I
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
S	S	L	L	L	L	L	L
Method	<u>Method</u>	Method	Method	Method	Method	Method	Method
9310	9310	EPA904.0	EPA 903.0	EPA 900.0	SM 7110C	EPA904.0	EPA 903.0
r Dr	1 <u>P</u> F	<u>DF</u>	<u>DF</u>	<u>DF</u> 1	<u>DF</u>	<u>DF</u>	1 <u>D</u> F
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
3	2	0.92	0.74	3.58	2.28	0.92	0.74
<u> </u>	<u>LOQ</u> 2	<u>LOQ</u> 1	<u>LOQ</u> 1	<u>LOQ</u>	<u>Гоо</u>	1	<u> 1</u>
Run Analyst	Run Analyst	Run Analyst	Run Analyst	Run Analyst	Run Analyst	Run Analyst	Run Analyst
05-May-11 MO	05-May-11 MO	02-May-11 MO	04-May-11 MO	30-Apr-11 MO	05-May-11 MO	02-May-11 MO	04-May-11 MO

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June 06, 2011

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	<u>Client ID#</u> RP-SED-03-110420	<u>Client ID#</u> RP-SED-03-110420	<u>Client ID#</u> RP-SED-02-110420	<u>Client ID#</u> RP-SED-02-110420	<u>Client ID#</u> RP-SED-02-110420	<u>Client ID#</u> RP-SED-02-110420	<u>Client ID#</u> RP-SED-01-110419	Client ID# RP-SED-01-110419
	<u>Lab ID#</u> 1107985-30	<u>Lab ID#</u> 1107985-30	<u>Lab ID#</u> 1107985-29	<u>Lab ID#</u> 1107985-29	<u>Lab ID#</u> 1107985-29	<u>Lab ID#</u> 1107985-29	<u>Lab ID#</u> 1107985-28	<u>Lab ID#</u> 1107985-28
	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11
	Analyte Gross Beta	<u>Analyte</u> Gross Alpha	<u>Analyte</u> Radium-228	<u>Analyte</u> Radium-226	Analyte Gross Beta	<u>Analyte</u> Gross Alpha	<u>Analyte</u> Radium-228	Analyte Radium-226
	Result 3.4 +/-0.1	Result 4.5 +/-0.16	<u>Result</u> U+/-0.47	Result 1.93 +/-0.49	Result 4.8 +/-0.1	<u>Result</u> 5.44 +/-0.18	Result U +/-0.34	Result 1.75 +/-0.47
1 1 0	<u>Units</u> pci/q	Units pci/g	<u>Units</u> pci/g	Units pci/g	Units pci/g	Units pci/g	Units pci/g	Units pci/g
	Matrix S	Matrix S	Matrix S	Matrix S	Matrix S	Matrix S	Matrix S	Matrix S
	<u>Method</u> 9310	<u>Method</u> 9310	<u>Method</u> 9320	<u>Method</u> 9315	<u>Method</u> 9310	<u>Method</u> 9310	<u>Method</u> 9320	<u>Method</u> 9315
	1 DF	DF	DF	DF	DF	DF 1	DF 1	DF 1
	3 <u>LOD</u>	<u>LOD</u> 2	LOD 2	LOD 1	<u>LOD</u>	<u>LOD</u> 2	<u>LOD</u> 2	1 00
	<u> LOQ</u>	<u>LOQ</u>	<u>LOQ</u> 2	1 1	<u>LOQ</u>	<u>LOQ</u> 2	<u>LOQ</u>	<u> </u>
	<u>Run Analyst</u> 06-May-11 MO	Run Analyst 06-May-11 MO	Run Analyst 04-May-11 MO	<u>Run</u> <u>Analyst</u> 04-May-11 MO	Run Analyst 06-May-11 MO	Run Analyst 06-May-11 MO	Run Analyst 04-May-11 MO	Run <u>Analyst</u> 04-May-11 MO

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Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

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						:	
<u>Client ID#</u> RP-SW-01-110419	<u>Client ID#</u> RP-SW-01-110419	Client ID# RP-SED-BG-041911	Client ID# RP-SED-BG-041911	Client ID# RP-SED-BG-041911	Client ID# RP-SED-BG-041911	<u>Client ID#</u> RP-SED-03-110420	RP-SED-03-110420
<u>Lab ID#</u> 1107985-32	<u>Lab ID#</u> 1107985-32	<u>Lab ID#</u> 1107985-31	<u>Lab ID#</u> 1107985-31	<u>Lab ID#</u> 1107985-31	<u>Lab ID#</u> 1107985-31	<u>Lab ID#</u> 1107985-30	1107985-30
Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 20-Apr-11	20-Apr-11
Analyte Gross Beta	Analyte Gross Alpha	Analyte Radium-228	<u>Analyte</u> Radium-226	Analyte Gross Beta	<u>Analyte</u> Gross Alpha	Analyte Radium-228	Radium-226
<u>Result</u> 28.1 +/-2.74	Result 75 +/-4.3	Result U +/-0.44	Result 1.71 +/-0.45	Result 1.34 +/-0.07	Result 1.9 +/-0.11	Result U +/-0.53	2.73 +/-0.55
Units pci/l	Units pci/I	Units pci/g	Units pci/g	<u>Units</u> pci/g	<u>Units</u> pci/g	<u>Units</u> pci/g	pci/g
Matrix L	Matrix L	Matrix S	Matrix S	Matrix S	Matrix S	Matrix S	S
Method EPA 900.0	<u>Method</u> SM 7110C	<u>Method</u> 9320	<u>Method</u> 9315	<u>Method</u> 9310	<u>Method</u> 9310	<u>Method</u> 9320	9315
DF.	<u>DF</u>	1 DF	DF 1	DF	DF	<u>DF</u>	<u> </u>
<u>LOD</u> 3.58	<u>LOD</u> 2.28	<u>LOD</u> 2	LOD 1	<u>LOD</u> 3	<u>LOD</u>	<u>LOD</u> 2	<u>ب</u> [2
<u>L00</u>	<u>LOQ</u> 3	<u>LOQ</u> 2	<u>LOQ</u> 1	<u>LOQ</u>	<u>LOQ</u> 2	<u>LOQ</u> 2	1 6
Run Analyst 30-Apr-11 MO	Run Analyst 05-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 06-May-11 MO	Run Analyst 06-May-11 MO	Run Analyst 04-May-11 MO	04-May-11 MO

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ENVIRONMENTAL TECHNOLOGIES, INC.

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

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Client ID#	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	Client ID#
SPOP35-110421	SPOP35-110421	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-01-110419	RP-SW-01-110419
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-34	1107985-34	1107985-33	1107985-33	1107985-33	1107985-33	1107985-32	1107985-32
Collected	Collected	Collected	Collected	Collected	Collected	Collected	Collected
21-Apr-11	21-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11
Analyte	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	Analyte	<u>Analyte</u>
Gross Beta	Gross Alpha	Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226
Result 2314.11 +/-73.39	<u>Result</u> 1326 +/-18	<u>Result</u> U +/-0.39	Result U +/-0.26	<u>Result</u> U +/-1.64	Result U +/-1,34	Result U +/-0.34	<u>Result</u> U +/-0.28
Units	<u>Units</u>	Units	Units	<u>Units</u>	<u>Units</u>	Units	Units
pci/l	pci/l	pci/l	pci/I	pci/l	pci/l	pci/l	pci/l
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L	L	L	L	L	L	L
Method	Method	Method	<u>Method</u>	<u>Method</u>	Method	Method	Method
EPA 900.0	SM 7110C	EPA904.0	EPA 903.0	EPA 900.0	SM 7110C	EPA904.0	EPA 903.0
1 DF	DE 1	1 <u>DF</u>	1 1	DE 1	DF 1	DF	DF 1
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
3.58	2.28	0.92	0.74	3.58	2.28	0.92	0.74
<u>Loa</u>	3	1	1	<u>LOQ</u>	<u>LOQ</u>	1 1	<u>LOQ</u> 1
Run Analyst 11-May-11 MO	Run Analyst 05-May-11 MO	Run Analyst 02-May-11 MO	Run <u>Analyst</u> 04-May-11 MO	Run Analyst 30-Apr-11 MO	Run Analyst 05-May-11 MO	Run Analyst 02-May-11 MO	Run Analyst 04-May-11 MO

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<u>Client ID#</u>	Client ID#	Client ID#	<u>Client ID#</u>	Client ID#	Client ID#
MW-RM-110420	MW-RM-110420	MW-RM-110420	MW-RM-110420	SPOP35-110421	SPOP35-110421
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-35	1107985-35	1107985-35	1107985-35	1107985-34	1107985-34
Collected	Collected	Collected	Collected	Collected	Collected
21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11
<u>Analyte</u>	Analyte	Analyte	Analyte	<u>Analyte</u>	<u>Analyte</u>
Radium-228	Radium-226	Gross Beta	Gross Alpha	Radium-228	Radium-226
Result U +/-0.27	<u>Result</u> U +/-0.29	Result 11.22 +/-2.95	Result U +/-1.2	Result 17.3 +/-1.76	Result 17.29 +/-1.27
Units	<u>Units</u>	Units	Units	Units	<u>Units</u>
pci/l	pci/l	pci/l	pci/l	pci/l	pci/l
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L	L		L	L
Method	Method	Method	Method	Method	<u>Method</u>
EPA904.0	EPA 903.0	EPA 900.0	SM 7110C	EPA904.0	EPA 903.0
<u>DF</u>	1 DE	1 DF	<u>DF</u>	<u>DF</u> 1	<u>DF</u>
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
0.92	0.74	3.58	2.28	0.92	0.74
1 1	1 1	<u>LOQ</u>	<u>LOQ</u>	1 LOQ	<u>LOQ</u>
Run Analyst 03-May-11 MO	Run Analyst 04-May-11 MO	Run Analyst 30-Apr-11 MO	Run Analyst 05-May-11 MO	Run Analyst 02-May-11 MO	Run Analyst 04-May-11 MO

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Run Analyst 02-May-11 DHC	<u>Loa</u>	LOD	1 <u>DF</u>	Method SM2310-B	Matrix L	<u>Units</u> mg/L	<u>Result</u> 470	Analyte Alkalinity	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-03	Client ID# MW-1-110419
Run Analyst 02-May-11 DHC	<u>LOQ</u> 20	<u>LOD</u> 20	<u>DF</u>	Method SM 4500- CO2D	Matrix L	<u>Units</u> mg/L	Result ND	Analyte Carbonate as CaCO3	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-02	<u>Client ID#</u> MD-SW-110420
Run Analyst 02-May-11 DHC	<u>LOQ</u>	<u>LOD</u>	DF	<u>ttrix Method</u> L SM-17 2320 B	Matrix L	<u>Units</u> mg/L	Result 317	<u>Analyte</u> Bicarbonate	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-02	<u>Client ID#</u> MD-SW-110420
Run Analyst 02-May-11 DHC	<u>LOQ</u>	LOD	DF	<u>Method</u> SM2310-B	Matrix L	Units mg/L	Result 322	<u>Analyte</u> Alkalinity	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-02	Client ID# MD-SW-110420
Run <u>Analyst</u> 02-May-11 DHC	<u>LOQ</u> 20	<u>LOD</u> 20	DF 1	Method SM 4500- CO2D	Matrix L	Units mg/L	Result ND	<u>Analyte</u> Carbonate as CaCO3	Collected 21-Apr-11	<u>Lab ID#</u> 1107985-01	<u>Client ID#</u> JPOP41S-110421
Run Analyst 02-May-11 DHC	5 <u>LOQ</u>	<u>LOD</u>	DF	<u>Itrix</u> <u>Method</u> L SM-17 2320 B	Matrix L	<u>Units</u> mg/L	Result 999	<u>Analyte</u> Bicarbonate	Collected 21-Apr-11	<u>Lab ID#</u> 1107985-01	Client ID# JPOP41S-110421
Run <u>Analyst</u> 02-May-11 DHC	<u> </u>	<u>LOD</u>	<u>DF</u> 1	Method SM2310-B	Matrix L	<u>Units</u> mg/L	Result 1000	<u>Analyte</u> Alkalinity	Collected 21-Apr-11	<u>Lab ID#</u> 1107985-01	<u>Client ID#</u> JPOP41S-110421

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Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

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<u>Client ID#</u> MW-1-110419
<u>Lab ID#</u> 1107985-03
Collected 19-Apr-11
<u>Analyte</u> Bicarbonate
<u>Result</u> 465
<u>Units</u> mg/L
Matrix L SM
<u>Method</u> SM-17 2320 B
DF 1
<u> LOD</u>
<u>L00</u>
Run Analyst 02-May-11 DHC

MW-1-110419	Client ID#	
1107985-03	Lab ID#	
19-Apr-11	Collected	
Carbonate as CaCO3	<u>Analyte</u>	
ND	Result	
mg/L	Units	
۲	Matrix	
SM 4500- CO2D	Method	
<u> </u>	DF	
20	LOD	
20	<u> </u>	
02-May-11 DHC	Run Analyst	

<u>Client ID#</u> MW-2-110421	Client ID# Lab ID# Collected Anal MW-2-110421 1107985-04 21-Apr-11 Alkalinit
<u>Lab ID#</u>	<u>Lab ID#</u> <u>Collected</u>
1107985-04	1107985-04 21-Apr-11
Collected 21-Apr-11	Collected 21-Apr-11
y <u>te</u>	<u>Analyte</u>
nate	Alkalinity
Result	<u>Result</u>
473	475
<u>Units</u>	<u>Units</u>
mg/L	mg/L
<u>Units</u>	<u>Units</u>
mg/L	mg/L
<u>Units</u>	<u>Units</u>
mg/L	mg/L
<u>Units</u>	<u>Units</u>
mg/L	mg/L
<u>Units</u>	<u>Units</u>
mg/L	mg/L
Units Matrix Method DF mg/L L SM-17 2320 1 B B	<u>Units</u> mg/L

	110,000	1107985-04	<u>Lab ID#</u>	
	1	21-Δnr-11	Collected	
	כמו סטו ומוכר מס כמכסט	Carbonate as CaCO3	Analyte	
	ē	5	Result	
		ma/l	Units	
	r		Matrix	
	CO2D	SM 4500-	Method	
	۲	-	DF	
***************************************	7.0	20	LOD	
	7	20	LOQ	
	or hay it bic	02-May-11 DHC	Run Analyst	

MW-2-110421

Client ID#

	MW-2-2-110421	Client ID# Lab ID# Collected Analyte	MW-2-2-110421	Client ID#
	1107985-05	Lab ID#	1107985-05	Lab ID#
	21-Apr-11	Collected	21-Apr-11	Collected
	Bicarbonate	<u>Analyte</u>	Alkalinity	<u>Analyte</u>
	398	Result	400	Result
	mg/L	Units	mg/L	<u>Units</u>
	L S	Matrix	_	Matrix
В	SM-17 2320	Method	SM2310-B	Method
	<u></u>	PF	—	DF
	5	LOD		LOD
	σ	<u>L00</u>	5	LOQ
	02-May-11 DHC	Matrix Method DF LOD LOQ Run Analyst	02-May-11 DHC	Run Analyst

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Run <u>Analyst</u> 02-May-11 DHC	<u>LOQ</u> 20	<u>LOD</u> 20	DF 1	Method SM 4500- CO2D	Matrix L	<u>Units</u> mg/L	Result ND	Analyte Carbonate as CaCO3	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-07	<u>Client ID#</u> MW-4-110420
Run Analyst 02-May-11 DHC	5 <u>LOQ</u>	5	ㅂ	Method SM-17 2320 B	Matrix L	<u>Units</u> mg/L	<u>Result</u> 450	<u>Analyte</u> Bicarbonate	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-07	Client ID# MW-4-110420
Run Analyst 02-May-11 DHC	<u>LOQ</u>	LOD	<u>DF</u>	Method SM2310-B	<u>Matrix</u> L	Units mg/L	Result 451	Analyte Alkalinity	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-07	Client ID# MW-4-110420
Run <u>Analyst</u> 02-May-11 DHC	<u>LOQ</u> 20	<u>LOD</u> 20	DE	Method SM 4500- CO2D	Matrix L	Units mg/L	Result ND	<u>Analyte</u> Carbonate as CaCO3	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-06	<u>Client ID#</u> MW-3-110420
Run Analyst 02-May-11 DHC	<u> </u>	<u>LOD</u>	DF	atrix <u>Method</u> L SM-17 2320 B	I≤	Units mg/L	Result 311	<u>Analyte</u> Bicarbonate	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-06	<u>Client ID#</u> MW-3-110420
Run Analyst 02-May-11 DHC	<u>LOQ</u>	LOD	<u>DF</u>	<u>Method</u> SM2310-B	Matrix L	<u>Units</u> mg/L	Result 312	<u>Analyte</u> Alkalinity	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-06	<u>Client ID#</u> MW-3-110420
Run Analyst 02-May-11 DHC	<u>LOQ</u> 20	<u>LOD</u> 20	<u>DF</u> 1	Method SM 4500- CO2D	Matrix L	Units mg/L	Result ND	Analyte Carbonate as CaCO3	<u>Collected</u> 21-Apr-11	<u>Lab ID#</u> 1107985-05	<u>Client ID#</u> MW-2-2-110421

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<u>Cli</u>	<u>CI</u>	<u>Cli</u>	<u>CI:</u>	MW.	MW CI:	MM CI
<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>
MW-8-110419	MW-7-110419	MW-7-110419	MW-7-110419	MW-6-110420	MW-6-110420	MW-6-110420
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-10	1107985-09	1107985-09	1107985-09	1107985-08	1107985-08	1107985-08
Collected	Collected	Collected	Collected	Collected	Collected	Collected
19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
Analyte	Analyte	<u>Analyte</u>	<u>Analyte</u>	Analyte Carbonate as CaCO3	Analyte	Analyte
Alkalinity	Carbonate as CaCO3	Bicarbonate	Alkalinity		Bicarbonate	Alkalinity
Result	Result	Result	Result	Result	Result	<u>Result</u>
452	ND	334	351	ND	369	370
Units	Units	Units	Units	Units	Units	<u>Units</u>
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L	L S	L	L	L S	L
Method SM2310-B	Method SM 4500- CO2D	<u>Method</u> SM-17 2320 B	Method SM2310-B	Method SM 4500- CO2D	<u>Method</u> SM-17 2320 B	Method SM2310-B
DF 1	1 DF	DF	1 1	DF 1	r DF	DF
<u>LOD</u>	<u>LOD</u> 20	<u>LOD</u>	<u>LOD</u>	<u>LOD</u> 20	<u> </u>	<u>LOD</u>
<u>LOQ</u>	<u>LOQ</u> 20	5	5 LOO	<u>LOQ</u> 20	5 5	<u>LOQ</u>
Run Analyst	Run Analyst	Run Analyst	Run Analyst	Run Analyst	Run Analyst	Run Analyst
02-May-11 DHC	02-May-11 DHC	02-May-11 DHC	02-May-11 DHC	02-May-11 DHC	02-May-11 DHC	02-May-11 DHC

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Client ID#

Lab ID#

Collected

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	Bicarbonate	Analyte
	445	Result
	mg/L	Units
	L SM-17	Matrix Method
В	2320	ethod
		DF
	Сī	LOD
	ហ	<u>L00</u>
	02-May-11 DHC	Run Analyst

MW-8-110419	Client ID#		MW-8-110419
1107985-10 19-Apr-11 Carbonate a	Lab ID#		1107985-10 19-Apr-11 Bicarbonate
19-Apr-11	Collected		19-Apr-11
as Ca	Analyte		Bicarbonate
CO3			
ND ND	Result		445
mg/L	Units		mg/L
F	Matrix		_
SM 4500- CO2D	Method		SM-17 2320 B
SM 4500- 1 CO2D	DF	C	SM-17 2320 1
SM 4500- 1 20 CO2D			SM-17 2320 1 5
H	DF		SM-17 2320 1 5 5

MW-MD-02-110420 1107985-11 20-Apr-11 Bicarbonate	Client ID# <u>Lab ID#</u> <u>Collected</u> <u>Analyte</u>	<u>Client ID#</u> <u>Lab ID#</u> <u>Collected</u> <u>Analyte</u> MW-MD-02-110420 1107985-11 20-Apr-11 Alkalinity
nate 567		lyte Result ty 570
mg/L	Units	<u>Units</u> mg/L
L SI	1	Matrix L
SM-17 2320	Method	x <u>Method DF LOD LOQ</u> SM2310-B 1 5
Ľ	DF	<u>DF</u>
5	LOD	<u> </u>
5	<u>L</u> 00	5 <u>TOO</u>
02-May-11 DHO	Run Analys	Run <u>Analyst</u> 02-May-11 DHC

-		_
MW-MD-02-110420 1107985-11 20-Apr-11 Carbonate as CaCO3	Client ID#	MW-MD-02-110420 1107985-11 20-Apr-11 Bicarbonate
₽	nt II	₹
02-1	#	02-1
110		110
120		120
<u></u>	<u></u>	
1079	Lab ID#	1079
385-	\mathcal{L}	985-
j		11
20	C	20
Αp	Collected	Ą
7-11	led.	71
_		
arb	l>	3ica
ona	Analyte	por.
fe a	/te	nate
s C		
Š		
w		
	Re	(17
ND	Result	567
3	_	3
mg/L	Units	mg/L
	Matrix	
	 X	
		SM-17 2320 B
, MS	Z	17
SM 4500- CO2D	Method	232(
0 4		вО
<u> </u>	DF	ь.
	_	
20	LOD	Q
	,_	
20	<u>100</u>	5
0	ă	
0		02-May-11 DHC
2-M	Ru	Ź-M
ay-1		ay-i
02-May-11 DHC	Run Analys	i i
ž	vst	X K

	MW-MD-110420	Client ID#	MW-MD-110420	Client ID#
	1107985-12	Lab ID#	1107985-12	<u>Lab ID#</u>
	20-Apr-11	Collected	20-Apr-11	Collected
	Bicarbonate	Analyte	Alkalinity	Analyte
	577	Result	580	Result
	mg/L	Units	mg/L	Units
	_	Matrix	_	Matrix
В	SM-17 2320	Method	SM2310-B	Method
	<u></u>	DF	<u>-</u>	DF
	ហ	TOD		<u>LOD</u>
		Ö		D
	И	<u>100</u>	5	<u>LOQ</u>

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Client ID#	<u>Client ID#</u>	<u>Client ID#</u>	Client ID#	Client ID#	Client ID#	<u>Client ID#</u>
PR-SW-01-110420	PR-SW-01-110420	PR-SW-01-110420	NPOP20E-110420	NPOP20E-110420	NPOP20E-110420	MW-MD-110420
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-15	1107985-15	1107985-15	1107985-13	1107985-13	1107985-13	1107985-12
Collected	Collected 20-Apr-11	Collected	Collected	Collected	Collected	Collected
20-Apr-11		20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
Analyte Carbonate as CaCO3	Analyte	<u>Analyte</u>	Analyte	<u>Analyte</u>	Analyte	Analyte
	Bicarbonate	Alkalinity	Carbonate as CaCO3	Bicarbonate	Alkalinity	Carbonate as CaCO3
Result ND	<u>Result</u> 75.0	Result 80	Result ND	Result 590	Result 590	Result ND
<u>Units</u>	Units	<u>Units</u>	<u>Units</u>	<u>Units</u>	<u>Units</u>	<u>Units</u>
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L S	L	L	L s	L	L
Method SM 4500- CO2D	<u>Method</u> SM-17 2320 B	Method SM2310-B	Method SM 4500- CO2D	<u>Method</u> SM-17 2320 B	Method SM2310-B	Method SM 4500- CO2D
DF 1	1 DF	<u>DF</u>	DF 1	<u>DF</u>	1 DE	DF 1
<u>LOD</u> 20	5 <u>LOD</u>	LOD	<u>LOD</u> 20	LOD 5	<u>LOD</u>	<u>LOD</u> 20
<u>LOQ</u> 20	<u>۲</u> ٥۵	<u>LOQ</u>	<u>LOQ</u> 20	<u>LOQ</u>	<u> 100</u>	<u>LOQ</u> 20
Run Analyst	Run Analyst	Run Analyst	Run <u>Analyst</u>	Rum Analyst	Run Analyst	Run <u>Analyst</u>
02-May-11 DHC	02-May-11 DHC	02-May-11 DHC	02-May-11 DHC	02-May-11 DHC	02-May-11 DHC	02-May-11 DHC

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RB-01-110420

<u>Lab ID#</u> 1107985-16

Collected 20-Apr-11

<u>Analyte</u> Alkalinity

Client ID#

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< 5.0	Result
mg/L	Units
_	Matrix
SM2310-B	Method
⊢	DF
	LOD
Vī	LOQ
02-May-11 DHC	Run Analyst

	RB-01-110420	Client ID#	<u>Client ID#</u> RB-01-110420
	1107985-16	Lab ID#	<u>Lab ID# Collected</u> £ 1107985-16 20-Apr-11 Bica
	20-Apr-11	Collected	Collected 20-Apr-11
	Carbonate as CaCO3	Analyte	Analyte Bicarbonate
	ND	Result	Result ND
	ND mg/L	Result Units	Result Units ND mg/L
			Units Matri
CO2D	mg/L	Units Matrix Method	Units Matri
CO2D	mg/L L SM 4500- 1	Units Matrix Method DF	Units Matri
CO2D	mg/L L SM 4500-	Units Matrix Method DF	Units Matri
CO2D	mg/L L SM 4500- 1	Units Matrix Method DF LOD LOQ	Units Matrix Method DF mg/L L SM-17 2320 1

<u>Client ID#</u> RB-02-110421	Client ID# RB-02-110421
<u>Lab ID#</u> 1107985-17	<u>Lab ID#</u> <u>Collected</u> <u>Analy</u> 1107985-17 21-Apr-11 Alkalinity
Collected 21-Apr-11	Collected 21-Apr-11
<u>Analyte</u> Bicarbonate	<u>Analyte</u> Alkalinity
Client ID# Lab ID# Collected Analyte RB-02-110421 1107985-17 21-Apr-11 Bicarbonate ND	Client ID# Lab ID# Collected Analyte Result RB-02-110421 1107985-17 21-Apr-11 Alkalinity < 5.0
<u>Units</u> mg/L	<u>Units</u> <u>Ma</u> mg/L
Matrix L	_ F 5 .
Method SM-17 2320 B	Method SM2310-B
<u>DF</u>	1 <u>DF</u>
LOD 5	TOD
<u>LOQ</u>	<u>5</u>
x Method DF LOD LOQ Run Analyst SM-17 2320 1 5 5 02-May-11 DHC B	x Method DF LOD LOQ Run Analyst SM2310-B 1 5 02-May-11 DHC

Client ID# RM-JM-SW-02	Client ID# RB-02-110421
Lab ID# Collected Analyt 1107985-20 20-Apr-11 Alkalinity	<u>Lab ID#</u> 1107985-17
Collected 4 20-Apr-11 Alka	Collected 21-Apr-11
<u>Analyte</u> Alkalinity	Analyte oonate as CaCO3
Result 290	Result ND
Result Units 290 mg/L	Result <u>Units</u> ND mg/L
<u>Units</u> <u>Ma</u> mg/L	<u>Units</u> <u>Ma</u> mg/L
<u>Units</u> <u>Ma</u> mg/L	<u>Units</u> <u>M</u> a
<u>Units</u> <u>Ma</u> mg/L	<u>Units</u> <u>M</u> a
<u>Units</u> <u>Ma</u> mg/L	<u>Units</u> <u>M</u> a
Units Matrix Method DF mg/L L SM2310-B 1	<u>Units</u> <u>M</u> a

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RM-SW-BG-110419	RM-SW-BG-110419	RM-SW-BG-110419	RM-JM-SW-110420	RM-JM-SW-110420	RM-JM-SW-110420	RM-JM-SW-02
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-23	1107985-23	1107985-23	1107985-21	1107985-21	1107985-21	1107985-20
Collected 19-Apr-11	Collected	Collected	Collected	Collected	Collected	Collected
	19-Apr-11	19-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
<u>Analyte</u>	<u>Analyte</u>	Analyte	Analyte Carbonate as CaCO3	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>
Carbonate as CaCO3	Bicarbonate	Alkalinity		Bicarbonate	Alkalinity	Bicarbonate
Result ND	Result 246	Result 250	Result ND	Result 287	Result 291	<u>Result</u> 286
<u>Units</u>	Units	Units	<u>Units</u>	Units	Units	<u>Units</u>
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	L S	L	L	L S	L	L S
Method SM 4500- CO2D	Method SM-17 2320 B	Method SM2310-B	Method SM 4500- CO2D	<u>Method</u> SM-17 2320 B	Method SM2310-B	Method SM-17 2320 B
<u>DF</u>	<u>DF</u>	1 PF	1 <u>DF</u>	1 DF	<u>DF</u>	1 <u>DF</u>
<u>LOD</u> 20	<u> 100</u>	<u>LOD</u>	<u>LOD</u> 20	5 <u>LOD</u>	<u>LOD</u>	5
<u>LOQ</u> 20	5 LOQ	<u>5</u>	<u>LOQ</u> 20	5 <u>LOO</u>	<u> 100</u>	5
Run Analyst	Run Analyst	Run Analyst	Run <u>Analyst</u>	Run <u>Analyst</u>	Run Analyst	Run Analyst
03-May-11 DHC	03-May-11 DHC	03-May-11 DHC	03-May-11 DHC	03-May-11 DHC	03-May-11 DHC	03-May-11 DHC

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<u>Client ID#</u> RP-SW-01-110419	<u>Client ID#</u> RP-JM-SW-110420	Client ID# RP-JM-SW-110420	<u>Client ID#</u> RP-JM-SW-110420	Client ID# RP-JM-SW-01- 110420	Client ID# RP-JM-SW-01- 110420	Client ID# RP-JM-SW-01- 110420
<u>Lab ID#</u>) 1107985-32	<u>Lab ID#</u>) 1107985-27	<u>Lab ID#</u>) 1107985-27	<u>Lab ID#</u> 0 1107985-27	<u>Lab ID#</u> 1107985-26	<u>Lab ID#</u> 1107985-26	<u>Lab ID#</u> 1107985-26
Collected 19-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11
<u>Analyte</u> Alkalinity	Analyte Carbonate as CaCO3	<u>Analyte</u> Bicarbonate	Analyte Alkalinity	Analyte Carbonate as CaCO3	<u>Analyte</u> Bicarbonate	<u>Analyte</u> Alkalinity
<u>Result</u> 333	<u>Result</u> ND	<u>Result</u> 282	Result 285	Result ND	<u>Result</u> 281	<u>Result</u> 284
Units mg/L	<u>Units</u> mg/L	Units mg/L	Units mg/L	<u>Units</u> mg/L	<u>Units</u> mg/L	<u>Units</u> mg/L
Matrix L	<u>Matrix</u> L	Matrix L S	Matrix L	Matrix L	Matrix L 2	Matrix L
<u>Method</u> SM2310-B	Method SM 4500- CO2D	ıtrix <u>Method</u> L SM-17 2320 B	Method SM2310-B	Method SM 4500- CO2D	atrix Method L SM-17 2320 B	Method SM2310-B
1 <u>P</u> F	1 <u>DF</u>	DF 1	DF 1	<u>DF</u>	1 F	<u>D</u> F
LOD	<u>LOD</u> 20	5 <u>TOD</u>	<u>LOD</u>	<u>LOD</u> 20	<u> LOD</u>	<u>LOD</u>
<u> </u>	<u>LOQ</u> 20	<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u> 20	5	<u>LOQ</u>
Run Analyst 03-May-11 DHC	Run <u>Analyst</u> 03-May-11 DHC	Run Analyst 03-May-11 DHC	Run Analyst 03-May-11 DHC	Run Analyst 03-May-11 DHC	Run <u>Analyst</u> 03-May-11 DHC	Run Analyst 03-May-11 DHC

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Client ID# Lab ID# SPOP35-110421 1107985-34	Client ID# Lab ID# SPOP35-110421 1107985-34	Client ID# Lab ID# SPOP35-110421 1107985-34	<u>Client ID#</u> RP-SW-BG-041911 1107985-33	<u>Client ID#</u> <u>Lab ID#</u> RP-SW-BG-041911 1107985-33	Client ID# Lab ID# RP-SW-BG-041911 1107985-33	
Collected 21-Apr-11	Collected 21-Apr-11	Collected 21-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	
Analyte Carbonate as CaCO3	<u>Analyte</u> Bicarbonate	<u>Analyte</u> Alkalinity	Analyte Carbonate as CaCO3	<u>Analyte</u> Bicarbonate	Analyte Alkalinity	
Result ND	Result 619	Result 620	Result ND	<u>Result</u> 339	<u>Result</u>	
Units mg/L	Units mg/L	Units mg/L	<u>Units</u> mg/L	<u>Units</u> mg/L	<u>Units</u> mg/L	
Matrix L	Matrix L	Matrix L	Matrix L	Matrix L	Matrix L	
Method SM 4500- CO2D	Method SM-17 2320 B	Method SM2310-B	Method SM 4500- CO2D	<u>Method</u> SM-17 2320 B	Method SM2310-B	
1 <u>D</u> F	1 DF	1 P	DF 1	DF	1 PF	
<u>LOD</u> 20	<u>LOD</u>	<u>[0</u>]	<u>LOD</u> 20	<u>LOD</u>	<u> </u>	
<u>LOQ</u> 20	2 <u>100</u>	<u> </u>	<u>LOQ</u> 20	5 <u>100</u>	<u>L00</u>	
Run Analyst 03-May-11 DHC	Run <u>Analyst</u> 03-May-11 DHC	Run Analyst 03-May-11 DHC	Run Analyst 03-May-11 DHC	Run Analyst 03-May-11 DHC	Run Analyst 03-May-11 DHC	

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Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

June 06, 2011

Analytical Laboratories

MW-RM-110420 1107985-35 Lab ID# Collected 21-Apr-11 Analyte Result 220

Client ID#

MW-RM-110420 Client ID# Client ID# 1107985-35 Lab ID# Lab ID# 21-Apr-11 Collected Collected Bicarbonate

MW-RM-110420

1107985-35 21-Apr-11 Carbonate as CaCO3 Alkalinity Analyte Analyte Result ND Result 218 <u>Units</u> mg/L <u>Units</u> mg/L Units mg/L Matrix Method
L SM-17 2320
B Matrix L Method SM2310-B SM 4500-CO2D Method <u>DF</u> DF. DF LOD 20 <u> LOD</u> 5 <u>100</u> <u> 5</u> 12-May-11 DHC 12-May-11 DHC 12-May-11 DHC Run Analyst Run Analyst Run Analyst

Fax: 330-253-4489

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ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

<u>Client ID#</u> MW-6-110420	Client ID# MW-4-110420	Client ID# MW-3-110420	Client ID# MW-2-2-110421	Client ID# MW-2-110421	Client ID# MW-1-110419	Client ID# MD-SW-110420	<u>Client ID#</u> JPOP41S-110421
<u>Lab ID#</u> 1107985-08	<u>Lab ID#</u> 1107985-07	<u>Lab ID#</u> 1107985-06	<u>Lab.ID#</u> 1107985-05	<u>Lab ID#</u> 1107985-04	<u>Lab ID#</u> 1107985-03	<u>Lab ID#</u> 1107985-02	<u>Lab ID#</u> 1107985-01
Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 21-Apr-11	Collected 21-Apr-11	Collected 19-Apr-11	Collected 20-Apr-11	Collected 21-Apr-11
Analyte Total Dissolved Solids (TDS)	<u>Analyte</u> Total Dissolved Solids (TDS)	Analyte Total Dissolved Solids (TDS)	Analyte Total Dissolved Solids (TDS)	Analyte Total Dissolved Solids (TDS)			
<u>Result</u> 2690	Result 1210	Result 785	Result 3330	<u>Result</u> 3500	Result 2000	<u>Result</u> 3860	<u>Result</u> 6370
Units mg/L	<u>Units</u> mg/L	<u>Units</u> mg/L	Units mg/L	<u>Units</u> mg/L	<u>Units</u> mg/L	Units mg/L	Units mg/L
Matrix L	Matrix L	Matrix L	Matrix L	Matrix L	Matrix L	Matrix L	Matrix L
Method SM 2540-C	<u>Method</u> SM 2540-C	Method SM 2540-C	Method SM 2540-C	Method SM 2540-C	Method SM 2540-C	Method SM 2540-C	Method SM 2540-C
1 PF	DF	DF 1	<u>DF</u>	<u>DF</u>	<u>DF</u>	1 DF	1 DF
F LOD	<u>LOD</u>	<u>LOD</u> 5	<u>LOD</u>	<u>LOD</u> 5	<u> 100</u>	<u>LOD</u>	<u>LOD</u>
5 <u>100</u>	5	<u>LOQ</u>	5	<u>Loa</u>	5	<u> 100</u>	<u>LOQ</u>
Run Analyst 29-Apr-11 JRK	Run Analyst 29-Apr-11 JRK	Run Analyst 29-Apr-11 JRK	Run Analyst 29-Apr-11 JRK				

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Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

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Client ID# RB-02-110421	Client ID# RB-01-110420	Client ID# PR-SW-01-110420	Client ID# NPOP20E-110420	<u>Client ID#</u> MW-MD-110420	Client ID# MW-MD-02-110420	<u>Client ID#</u> MW-8-110419	<u>Client ID#</u> MW-7-110419
<u>Lab ID#</u> 1107985-17	<u>Lab ID#</u> 1107985-16	<u>Lab ID#</u> 1107985-15	<u>Lab ID#</u> 1107985-13	<u>Lab ID#</u> 1107985-12	<u>Lab ID#</u> 1107985-11	<u>Lab ID#</u> 1107985-10	<u>Lab ID#</u> 1107985-09
Collected 21-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11
Analyte Total Dissolved Solids (TDS)							
Result 60	Result 100	Result 7950	Result 4440	<u>Result</u> 7250	<u>Result</u> 5490	Result 940	Result 515
Units mg/L	<u>Units</u> mg/L	Units mg/L	<u>Units</u> mg/L	<u>Units</u> mg/L	Units mg/L	<u>Units</u> mg/L	<u>Units</u> mg/L
Matrix L	Matrix L	Matrix L	Matrix L	<u>Matrix</u> L	Matrix L	Matrix L	Matrix L
Method SM 2540-C	<u>Method</u> SM 2540-C	Method SM 2540-C	Method SM 2540-C	Method SM 2540-C	<u>Method</u> SM 2540-C	Method SM 2540-C	<u>Method</u> SM 2540-C
DF 1	<u>DF</u>	DF 1	DF	r D	<u>DF</u>	<u>DF</u>	<u>DF</u>
5	5	<u>LOD</u>	5 <u>LOD</u>	5 100	<u> LOD</u>	5 <u>FOD</u>	<u>LOD</u>
<u> 100</u>	<u> 100</u>	<u>100</u>	<u>100</u>	<u> </u>	5 LOQ	5 <u>100</u>	<u> 100</u>
Run Analyst 29-Apr-11 JRK							

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29-Apr-11 JRK	5	5	1	SM 2540-C	_	mg/L	9360	Total Dissolved Solids (TDS)	21-Apr-11	1107985-34	SPOP35-110421	
Run	100	LOD	DF	Method	Matrix	Units	Result	Analyte	Collected	Lab ID#	Client ID#	
Run Analysı 29-Apr-11 JRK	2 <u>100</u>	2 <u>TOD</u>	1 PF	Method SM 2540-C	<u>Matrix</u> L	<u>Units</u> mg/L	Result 420	Analyte Total Dissolved Solids (TDS)	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-33	Client ID# RP-SW-BG-041911	
Run Analyst 29-Apr-11 JRK	<u>100</u>	<u> 100</u>	1 <u>P</u> F	<u>Method</u> SM 2540-C	Matrix L	<u>Units</u> mg/L	Result 2120	Analyte Total Dissolved Solids (TDS)	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-32	Client ID# RP-SW-01-110419	
Run <u>Analyst</u> 29-Apr-11 JRK	<u> </u>	<u>LOD</u>	<u>DF</u>	<u>Method</u> SM 2540-C	Matrix L	Units mg/L	Result 1220	Analyte Total Dissolved Solids (TDS)	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-27	Client ID# RP-JM-SW-110420	
Run Analysi 29-Apr-11 JRK	<u> </u>	LOD 5	<u>DF</u>	Method SM 2540-C	Matrix L	<u>Units</u> mg/L	Result 1800	Analyte Total Dissolved Solids (TDS)	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-26	Client ID# RP-JM-SW-01- 110420	
Run Analyst 29-Apr-11 JRK	<u>1500</u>	<u> 100</u>	DF	<u>Method</u> SM 2540-C	Matrix L	<u>Units</u> mg/L	Result 1260	Analyte Total Dissolved Solids (TDS)	Collected 19-Apr-11	<u>Lab ID#</u> 1107985-23	Client ID# RM-SW-BG-110419	
Run Analys 29-Apr-11 JRK	<u> </u>	2 <u>100</u>	1 DF	<u>Method</u> SM 2540-C	<u>Matrix</u> L	<u>Units</u> mg/L	Result 2410	Analyte Total Dissolved Solids (TDS)	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-21	Client ID# RM-JM-SW-110420	
Run Analysi 29-Apr-11 JRK	<u> 100</u>	<u>LOD</u>	<u>DF</u>	Method SM 2540-C	Matrix L	Units mg/L	<u>Result</u> 1920	Analyte Total Dissolved Solids (TDS)	Collected 20-Apr-11	<u>Lab ID#</u> 1107985-20	Client ID# RM-JM-SW-02	

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3310 Win Street Cuyahoga Falls, Ohio 44223

ENVIRONMENTAL TECHNOLOGIES, INC.

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MW-RM-110420 Client ID#

Lab ID#

1107985-35

21-Apr-11

Collected

Analyte

Total Dissolved Solids (TDS)

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

Dallas, TX 75354

Project #: Jackpile Received: 4/26/2011

<u>Result</u> 1340

mg/L Units

Matrix L

 $\frac{\text{Method}}{\text{SM 2540-C}} \quad \frac{\text{DF}}{1}$

<u>LOD</u>

<u> 100</u>

Run <u>Analyst</u> 12-May-11 JRK

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

<u>Client ID#</u>	Client ID#	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>
MW-6-110420	MW-4-110420	MW-3-110420	MW-2-2-110421	MW-2-110421	MW-1-110419	MD-SW-110420	JPOP41S-110421
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
110 7 985-08	1107985-07	1107985-06	1107985-05	1107985-04	1107985-03	1107985-02	1107985-01
Collected	Collected	Collected	Collected	Collected	Collected	Collected	Collected
20-Apr-11	20-Apr-11	20-Apr-11	21-Apr-11	21-Apr-11	19-Apr-11	20-Apr-11	21-Apr-11
<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	Analyte	<u>Analyte</u>	<u>Analyte</u>	Analyte
Mercury	Mercury	Mercury	Mercury	Mercury	Mercury	Mercury	Mercury
<u>Flag</u> U	Flag	Flag	Flag	Flag	Flag U	Flag U	Flag
<u>Result</u> <0.00006	Result < 0.00006	Result < 0.00006	Result < 0.00006	Result < 0.00006	Result < 0.00006	<u>Result</u> <0.00006	<u>Result</u> <0.00006
<u>Units</u>	<u>Units</u>	<u>Units</u>	<u>Units</u>	Units	Units	<u>Units</u>	Units
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L
Matrix L	Matrix L	Matrix L	<u>Matrix</u> L	<u>Matrix</u>	<u>Matrix</u>	<u>Matrix</u> L	Matrix L
Method	<u>Method</u>	<u>Method</u>	Method	<u>Method</u>	<u>Method</u>	Method	<u>Method</u>
7470A	7470A	7470A	7470A	7470A	7470A	7470A	7470A
<u>DF</u> <u>L(</u> 1 0.00	<u>DF LC</u> 1 0.00	<u>DF</u> L0	<u>DF</u> <u>L</u> (1 0.00	<u>DF LOD</u> 1 0.0000	<u>DF</u> <u>L</u> !	DF L	<u>DF</u> L 1 0.0
<u>DF LOD LOQ Run Analyst</u>	<u>LOD</u> <u>LOQ</u>	DF LOD LOQ Rum Analyst 1 0.00006 0.00020 04-May-11 VVK	<u>DF LOD LOQ</u>	<u>LOD</u> <u>LOQ</u>	<u>DF LOD LOQ Run Analyst</u>	<u>DF LOD LOQ Run Analyst</u>	<u>LOD</u> <u>LOQ</u>
1 0.00006 0.00020 04-May-11 VVK	0.00006 0.00020		1 0.00006 0.00020	0.00006 0.00020	1 0.00006 0.00020 04-May-11 VVK	1 0.00006 0.00020 04-May-11 VVK	0.00006 0.00020
<u>) Q</u> 0020 0	<u>)Q</u> 0020 0	<u>LOQ</u>).00020 0			<u>LOQ</u>).00020 0	<u>LOQ</u>).00020 C	
<u>Run</u> <u>/</u>	Run <u>Analyst</u>	Run Analyst	Run Analyst	Run <u>Analyst</u>	<u>Run</u>	<u>Run</u>	Run Analyst
)4-May-1	04-May-11 VVK)4-May-11 VVK	04-May-11 VVK	04-May-11 VVK)4-May-1)4-May-1	04-May-11 VVK
Analyst 11 VVK	Analyst -11 VVK	Analyst 1 VVK	Analyst 1 VVK	Analyst 11 VVK	Analyst 11 VVK	Analyst 11 VVK	Analyst -11 VVK

"Analytical Integrity"

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ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

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	<u>Client ID#</u> RB-01-110420	Client ID# PR-SW-01-110420	<u>Client ID#</u> PR-SED-01-110420	<u>Client ID#</u> NPOP20E-110420	<u>Client ID#</u> MW-MD-110420	Client ID# MW-MD-02-110420	<u>Client ID#</u> MW-8-110419	<u>Client ID#</u> MW-7-110419
		.00	20	D		20		
	<u>Lab ID#</u> 1107985-16	<u>Lab ID#</u> 1107985-15	<u>Lab ID#</u> 1107985-14	<u>Lab ID#</u> 1107985-13	<u>Lab ID#</u> 1107985-12	<u>Lab ID#</u> 1107985-11	<u>Lab ID#</u> 1107985-10	<u>Lab ID#</u> 1107985-09
	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11
	Analyte Mercury	<u>Analyte</u> Mercury	<u>Analyte</u> Mercury	<u>Analyte</u> Mercury	<u>Analyte</u> Mercury	Analyte Mercury	<u>Analyte</u> Mercury	<u>Analyte</u> Mercury
	Flag U	Flag U	Flag	Flag	Flag U	Flag U	Flag U	Flag
	<u>Result</u> <0.00006	Result < 0.00006	Result <0.0632	Result < 0.00006	Result < 0.00006	<u>Result</u> <0.00006	Result < 0.00006	Result < 0.00006
	<u>Units</u> mg/L	<u>Units</u> mg/L	Units mg/Kg	Units mg/L	Units mg/L	<u>Units</u> mg/L	Units mg/L	Units mg/L
	Matrix L	Matrix L	Matrix S	Matrix L	Matrix L	Matrix L	Matrix L	Matrix L
	<u>Method</u> 7470A	Method 7470A	<u>Method</u> 7471A	<u>Method</u> 7470A	<u>Method</u> 7470A	<u>Method</u> 7470A	<u>Method</u> 7470A	<u>Method</u> 7470A
	<u>DF LOD</u> 1 0.00000	<u>DF LOD</u> 1 0.00006	<u>DF LOD</u> 1 0.0632	<u>DF</u> <u>LOD</u> 1 0.00000	<u>DF</u> <u>LOD</u> 1 0.00000	<u>DF LOD</u> 1 0.0000	<u>DF</u> <u>LOD</u> 1 0.00000	<u>DF</u> <u>LOD</u> 1 0.00000
	<u>LOD</u> <u>LOQ</u> 0.00006 0.00020	<u>LOD</u> <u>LOQ</u> 0.00006 0.00020	<u> </u>	<u>LOD</u> <u>LOQ</u> 0.00006 0.00020	<u>LOD</u> <u>LOQ</u> 0.00006 0.00020	<u>LOD</u> <u>LOQ</u> 0.00006 0.00020	LOD LOQ Run Analyst 0.00006 0.00020 04-May-11 VVK	<u>LOD</u> <u>LOQ</u> 0.00006 0.00020
1	<u>Run</u> 04-May	ő	Run Analyst 04-May-11 VVK	<u>Run</u> 04-May	Q Q	<u>Run</u> 04-May-	<u>Run</u> 04-May-	5
	Analyst -11 VVK	Run Analyst H-May-11 VVK	Analyst -11 WK	Analyst 11 WK	Run Analyst I-May-11 VVK	Analyst 11 WK	Analyst 11 WK	Run Analyst I-May-11 VVK

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ENVIRONMENTAL TECHNOLOGIES, INC.

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June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

	RP □	RM C	RM C	C RM	RM C	R™	R™	
	Client ID# RP-JM-SED-01	Client ID# RM-SW-BG-110419	Client ID# RM-SED-BG-110419	<u>Client ID#</u> RM-JM-SW-110420	<u>Client ID#</u> RM-JM-SW-02	Client ID# RM-JM-SED-110420	Client ID# RM-JM-SED-02-110420	<u>Client ID#</u> RB-02-110421
	<u>Lab ID#</u> 1107985-24	<u>Lab ID#</u> 1107985-23	<u>Lab ID#</u> 1107985-22	<u>Lab ID#</u> 1107985-21	<u>Lab ID#</u> 1107985-20	<u>Lab ID#</u> 1107985-19	<u>Lab ID#</u> 1107985-18	<u>Lab ID#</u> 1107985-17
	Collected 20-Apr-11	Collected 19-Apr-11	Collected 19-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 20-Apr-11	Collected 21-Apr-11
	<u>Analyte</u> Mercury	<u>Analyte</u> Mercury	<u>Analyte</u> Mercury	<u>Analyte</u> Mercury	Analyte Mercury	<u>Analyte</u> Mercury	<u>Analyte</u> Mercury	<u>Analyte</u> Mercury
	Flag U	Flag U	Flag	Flag U	Flag	Flag U	<u>Flag</u> U	Flag
	Result < 0.0468	Result < 0.00006	Result < 0.0563	<u>Result</u> <0.00006	<u>Result</u> <0.00006	Result < 0.0474	<u>Result</u> <0.0436	<u>Result</u> < 0.00006
	<u>Units</u> mg/Kg	Units mg/L	<u>Units</u> mg/Kg	<u>Units</u> mg/L	<u>Units</u> mg/L	Units mg/Kg	Units mg/Kg	<u>Units</u> mg/L
	Matrix S	Matrix L	Matrix S	Matrix L	Matrix L	Matrix S	Matrix S	Matrix L
	Method 7471A	Method 7470A	<u>Method</u> 7471A	<u>Method</u> 7470A	Method 7470A	<u>Method</u> 7471A	<u>Method</u> 7471A	<u>Method</u> 7470A
	<u>DF</u>	<u>DF</u>	DF DF	<u>DF</u>	DF 1 (<u>DF</u>	1 <u>DF</u>	DF 1 (
	<u>LOD</u> 0.0468	<u>LOD</u>	<u>LOD</u> 0.0563	<u>LOD</u> 0.00006	<u>LOD</u> 0.00006	<u>LOD</u> 0.0474	<u>LOD</u> 0.0436	<u>LOD</u> 0.00006
	<u>LOQ</u> 0.234	<u>LOD</u> <u>LOQ</u> 0.00006 0.00020	<u>LOQ</u> 0.282	<u>LOD</u> <u>LOQ</u> 0.00006 0.00020	<u>LOQ</u> 0.00020	<u>LOQ</u> 0.237	<u>LOQ</u> 0.218	<u>DF LOD LOQ</u> 1 0.00006 0.00020
Page 74	Run Analyst 04-May-11 VVK	Run Analyst 04-May-11 VVK	Run Analyst 04-May-11 VVK	Run Analyst 04-May-11 VVK	LOD LOQ Run Analyst 0.00006 0.00020 04-May-11 VVK	Run Analyst 04-May-11 VVK	Run Analyst 04-May-11 VVK	Run Analyst 04-May-11 VVK

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<u>Client ID#</u>	Client ID#	<u>Client ID#</u>	<u>Client ID#</u>	<u>Client ID#</u>	Client ID#	Client ID#	Client ID#
RP-SW-01-110419	RP-SED-BG-041911	RP-SED-03-110420	RP-SED-02-110420	RP-SED-01-110419	RP-JM-SW-110420	RP-JM-SW-01-110420	RP-JM-SED-110420
110419	±	±	<u>#</u>	±	±	±	<u>#</u>
	6-041911	-110420	!-110420	110419	-110420	-01-110√)-11042(
				-		120	
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-32	1107985-31	1107985-30	1107985-29	1107985-28	1107985-27	1107985-26	1107985
#	<u>⊯</u>	<u>⊯</u>	<u>)#</u>	<u>)#</u>	<u>⊯</u>	<u>)#</u>	<u>Lab ID#</u>
85-32	85-31	85-30	85-29	85-28	85-27	85-26	1107985-25
Collected	Collected	<u>Collected</u>	Collected	Collected	Collected	Collected	Collected
19-Apr-11	19-Apr-11	20-Apr-11	20-Apr-11	19-Apr-11	20-Apr-11	20-Apr-11	20-Apr-11
Analyte	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>	Analyte	Analyte	<u>Analyte</u>
Mercury	Mercury	Mercury	Mercury	Mercury	Mercury	Mercury	Mercury
169	10	. 16	IO.	, I G	. 10	i le	lo : :
Flag ∪	Flag U	Flag	Flag U	Flag ∪	Flag	Flag U	Flag
<u>Result</u> <0.00006	<u>Result</u> <0.0532	<u>Result</u> <0.0419	<u>Result</u> < 0.0515	<u>Result</u> <0.0511	<u>Result</u> <0.00006	<u>Result</u> <0.00006	Result < 0.0525
-						_	
<u>Units</u>	<u>Units</u>	<u>Units</u>	Units	Units	<u>Units</u>	Units	<u>Units</u>
mg/L	mg/Kg	mg/Kg	mg/Kg	mg/Kg	mg/L	mg/L	mg/Kg
Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix	Matrix
L	S	S	S	S	L	L	S
<u>Method</u>	<u>Method</u>	<u>Method</u>	<u>Method</u>	<u>Method</u>	Method	Method	<u>Method</u>
7470A	7471A	7471A	7471A	7471A	7470A	7470A	7471A
1 <u>DF</u>	<u>DF</u>	DF 1	DF 1	DF	1 DE	<u>DF</u>	1 <u>DF</u>
<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>	<u>LOD</u>
0.00006	0.0532	0.0419	0.0515	0.0511		0.00006	0.0525
<u>LOD</u> <u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u>	LOD LOQ Run Analyst	<u>LOD</u> <u>LOQ</u>	<u>LOQ</u>
0.00006 0.00020	0.266	0.209	0.258	0.255	0.00006 0.00020 04-May-11 VVK	0.00006 0.00020	0.263
	<u>Run</u> 04-May	<u>Run</u> 04-May	<u>Run</u> 04-May	<u>Run</u> 04-May	<u>Run</u> 04-May		<u>Run</u> 04-May
Run Analysi	Run Analyst	Run Analyst	Run Analyst	Run Analyst	Analyst	Run Analyst	Run Analyst
04-May-11 VVK	04-May-11 VVK	04-May-11 VVK	04-May-11 VVK	04-May-11 VVK	-11 VVK	04-May-11 VVK	04-May-11 VVK

"Analytical Integrity" 3310 Win Street Cuyahoga Falls, Ohio 44223

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NELAP Certified

Page 75

Web Site: www.settek.com

Analytical Laboratories

June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850

Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

Client ID# Lab ID# Collected Analy MW-RM-110420 1107985-35 21-Apr-11 Mercury	<u>Client ID#</u> SPOP35-110421	Client ID# RP-SW-BG-041911
<u>Lab ID#</u>	<u>Lab ID#</u>	<u>Lab ID#</u>
1107985-35	1107985-34	1107985-33
Collected	Collected	Collected
21-Apr-11	21-Apr-11	19-Apr-11
<u>Analyte</u>	<u>Analyte</u>	<u>Analyte</u>
Mercury	Mercury	Mercury
<u>Flag</u>	Flaq	Elag
U	U	U
Result <0.00006	Result < 0.00006	<u>Result</u> <0.00006
Units	Units	<u>Units</u>
mg/L	mg/L	mg/L
<u>Matrix</u>	Matrix	<u>Matrix</u>
L	L	L
<u>Method</u>	Method	<u>Method</u>
7470A	7470A	7470A
<u>DF</u>	1 <u>DF</u>	1 <u>DF</u>
<u>LOD</u>	<u>LOD</u> 0.00006	<u>LOD</u> 0.00006
<u>LOQ</u>	<u>LOQ</u>	<u>LOQ</u>
0.00020	0.00020	0.00020
<u>DF LOD LOQ Run Analyst</u>	<u>DF LOD LOQ Run Analyst</u>	<u>DF LOD LOQ Run Analyst</u>
1 0.00006 0.00020 04-May-11 WK	1 0.00006 0.00020 04-May-11 VVK	1 0.00006 0.00020 04-May-11 WK

Fax: 330-253-4489

Analytical Laboratories

Client: Weston Solutions
Address: 14160 Dallas Parkway, Suite 850
Dallas, TX 75354
Received: 4/26/2011

Project #: Jackpile

MW-4-110420	MW-4-110420	MW-4-110420	MW-4-110420	MW-3-110420	MW-3-110420	MW-3-110420	MW-3-110420	MW-2-2-110421	MW-2-2-110421	MW-2-2-110421	MW-2-2-110421	MW-2-110421	MW-2-110421	MW-2-110421	MW-2-110421	MW-1-110419	MW-1-110419	MW-1-110419	MW-1-110419	MD-SW-110420	MD-SW-110420	MD-SW-110420	MD-SW-110420	JPOP41S-110421	JPOP41S-110421	JPOP41S-110421	JPOP41S-110421	Client ID#
1107985-07	1107985-07	1107985-07	1107985-07	1107985-06	1107985-06	1107985-06	1107985-06	1107985-05	1107985-05	1107985-05	1107985-05	1107985-04	1107985-04	1107985-04	1107985-04	1107985-03	1107985-03	1107985-03	1107985-03	1107985-02	1107985-02	1107985-02	1107985-02	1107985-01	1107985-01	1107985-01	1107985-01	Lab ID#
Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Analyte
1560	0.115	0.165	14.9	381	0.115	0.165	7.90	2030	0.115	0.165	45.3	2460	0.11	0.165	39.7	1190	2.40	0.165	15.6	2050	0.115	0.165	363	3140	54.4	0.165	15.4	Result
U	5	5 _	•		5 _	5	Ŭ	U	5	5	<u></u>		5 _	5 U	,	J	Ŏ	5 U	0.	U	5	5 _		J		5 _		
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	Units
0.200	0.115	0.165	0.200	0.200	0.115	0.165	0.200		0.115	0.165	0.200				0.200				0.200		0.115	0.165	0.200	0.200	0.115	0.165	0.200	<u>LOD</u>
2.5	0.6	0.8	2.5	2.5	0.6	0.8	2.5	2.5	0.6	0.8				0.8							0.6			2.5	0.6	0.8	2.5	<u>L00</u>
<u>ш</u>		<u>,_</u>	<u> </u>	_	₽		<u>—</u>		<u>—</u>	j 4	<u>ш</u>	 4	⊢	Н	j _{ere} sk		—	-		<u> </u>		<u> </u>	⊢	<u> </u>		⊢	ш	DF
_	_	_		_	_	_	г—	_		-	_		_	_	_	_	_	_	_	_	_	_	_	_		_	<u>-</u>	Matrix
300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	x Method
KMG	KMG	KMG	KMG	KMG	KMG	ΚMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	l>
03-May-11 20-Apr-11	03-May-11 20-Apr-11	03-May-11 20-Apr-11	03-May-11 20-Apr-11	03-May-11 20-Apr-1	03-May-11 20-Apr-11	03-May-11 20-Apr-11	03-May-11 20-Apr-11	03-May-11 21-Apr-11	03-May-11 21-Apr-11	03-May-11 21-A _l	03-May-11 21-Apr-1:	03-May-11 21-Apr-11	03-May-11 21-Apr-11	03-May-11 21-Apr-11	03-May-11 21-Apr-11	03-May-11 19-Apr-11	03-May-11 19-Apr-11	03-May-11 19-Apr-11	03-May-11 19-Apr-1	03-May-11 20-Apr-11	03-May-11 20-Apr-11	03-May-11 20-Apr-11	03-May-11 20-Apr-1	03-May-11 21-Apr-11	03-May-11 21-A _l	03-May-11 21-Apr-11	03-May-11 21-Apr-11	Analyzed Coll
pr-11	pr-11	pr-11	pr-11	pr-11	pr-11	pr-11	pr-11	pr-11	pr-11	21-Apr-11	pr-11	pr-11	pr-11	pr-11	pr-11	pr-11	pr-11	21-Apr-11	pr-11	pr-11	Collected							

Phone: 330-253-8211

Fax: 330-253-4489

Analytical Laboratories

Client: Weston Solutions June 06, 2011

Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

PR-SW-01-110420 PR-SW-01-110420	PR-SW-01-110420	PR-SW-01-110420	NPOP20E-110420	NPOP20E-110420	NPOP20E-110420	NPOP20E-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-110420	MW-MD-02-110420	MW-MD-02-110420	MW-MD-02-110420	MW-MD-02-110420	MW-8-110419	MW-8-110419	MW-8-110419	MW-8-110419	MW-7-110419	MW-7-110419	MW-7-110419	MW-7-110419	MW-6-110420	MW-6-110420	MW-6-110420	MW-6-110420	Client ID#
1107985-15 1107985-15	1107985-15	1107985-15	1107985-13	1107985-13	1107985-13	1107985-13	1107985-12	1107985-12	1107985-12	1107985-12	1107985-11	1107985-11	1107985-11	1107985-11	1107985-10	1107985-10	1107985-10	1107985-10	1107985-09	1107985-09	1107985-09	1107985-09	1107985-08	1107985-08	1107985-08	1107985-08	Lab ID#
Nitrate Sulfate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Analyte
0.115 5330	0.165	145	3580	0.115	0.165	18.9	3790	0.115	0.165	128	3670	0.115	0.165	133	350	0.115	0.165	19.4	106	0.115	0.165	3.50	1650	0.115	0.165	32.9	Result
<u> </u>	<u> </u>			<u> </u>	<u> </u>			<u> </u>	<u>_</u>			<u>_</u>	<u> </u>			<u>_</u>	<u> </u>			<u>_</u>	<u>_</u>			<u>_</u>	<u>_</u>		io Qi
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	Units
0.115 0.200	0.165	0.200	0.200	0.115		0.200				0.200				0.200			0.165	0.200	0.200	0.115	0.165	0.200	0.200	0.115	0.165	0.200	LOD
2.5	3.0	2.5		0.6		2.5		0.6																0.6	0.8	2.5	LOQ
о, о. н н	н	ш.	<u></u>	<u></u>	<u>-</u>	<u></u>	<u>-</u>	1	<u></u>	<u></u>	<u></u>	<u></u>	<u>⊢</u>	j. □	1	1	1	1	1	1	1	1	1	1	3 1		
	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	Matrix
300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	x Method
KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	Analysi
03-May-11 20 03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 19	03-May-11 19	03-May-11 19	03-May-11	03-May-11 19	03-May-11 19	03-May-11 19	03-May-11 19	03-May-11 20	03-May-11 20	03-May-11 20	03-May-11 20	Analyzed
20-Apr-11 20-Apr-11	20-Apr-1	20-Apr-1	20-Apr-1	20-Apr-1	20-Apr-1	20-Apr-1	20-Apr-11	20-Apr-11	20-Apr-11	20-Apr-1:	20-Apr-11	20-Apr-1:	20-Apr-11	20-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-1	19-Apr-1	19-Apr-1	20-Apr-1	20-Apr-1	20-Apr-1:	20-Apr-1	Collected

Phone: 330-253-8211

June 06, 2011

ENVIRONMENTAL TECHNOLOGIES, INC.

Analytical Laboratories

Received: 4/26/2011 Client: Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354 Weston Solutions

Project #: Jackpile

RP-JM-SW-110420	RP-JM-SW-110420	RP-JM-SW-110420	RP-JM-SW-110420	RP-JM-SW-01-110420	RP-JM-SW-01-110420	RP-JM-SW-01-110420	RP-JM-SW-01-110420	RM-SW-BG-110419	RM-SW-BG-110419	RM-SW-BG-110419	RM-SW-BG-110419	RM-JM-SW-110420	RM-JM-SW-110420	RM-JM-SW-110420	RM-JM-SW-110420	RM-JM-SW-02	RM-JM-SW-02	RM-JM-SW-02	RM-JM-SW-02	RB-02-110421	RB-02-110421	RB-02-110421	RB-02-110421	RB-01-110420	RB-01-110420	RB-01-110420	RB-01-110420	Client ID#
1107985-27	1107985-27	1107985-27	1107985-27	1107985-26	1107985-26	1107985-26	1107985-26	1107985-23	1107985-23	1107985-23	1107985-23	1107985-21	1107985-21	1107985-21	1107985-21	1107985-20	1107985-20	1107985-20	1107985-20	1107985-17	1107985-17	1107985-17	1107985-17	1107985-16	1107985-16	1107985-16	1107985-16	Lab ID#
Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Analyte
642	0.115 U	0.165 U	13.0	1040	0.115 U	0.165 U	19.1	877	0.115 U	0.165 U	13.4	1920	0.115 U	0.165 U	24.5	1900	0.115 U	0.165 U	26.7	0.200 U	0.115 U	0.165 U	0.200 U	0.200 U	0.115 U	0.165 U	0.200 U	Result Q
mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	mg/L	Units
0.200	0.115	0.165	0.200	0.200	0.115	0.165	0.200	0.200	0.115	0.165	0.200	0.200	0.115	0.165	0.200	0.200	0.115	0.165	0.200	0.200	0.115	0.165	0.200	0.200	0.115	0.165	0.200	LOD
2.5	0.6	0.8	2.5	2.5	0.6	0.8	2.5	2.5	0.6	0.8	2.5	2.5	0.6	0.8	2.5	2.5	0.6	0.8	2.5	2.5	0.6	0.8	2.5	2.5			2.5	LOQ
<u> </u>	<u> </u>	H	<u> </u>		-	<u></u>	1	<u>ш</u>	_	<u>—</u>	<u>ш</u>	⊣	<u> </u>	<u> </u>	↦	_	<u></u>		<u></u>	<u>-</u>	<u>-</u>	<u></u>	-	1	1	_	j k	<u>DF</u>
_	_	_	_	_	_	_	г	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	Г	_	_	_	Matrix
300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	Method
KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	SMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	Analyst
03-May-11			03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-1	03-May-11	03-May-11	t Analyzed

Phone: 330-253-8211

Analytical Laboratories

Client: June 06, 2011

Client: Weston Solutions Address: 14160 Dallas Parkway, Suite 850 Dallas, TX 75354

Received: 4/26/2011 Project #: Jackpile

MW-RM-110420	MW-RM-110420	MW-RM-110420	MW-RM-110420	SPOP35-110421	SPOP35-110421	SPOP35-110421	SPOP35-110421	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-BG-041911	RP-SW-01-110419	RP-SW-01-110419	RP-SW-01-110419	RP-SW-01-110419	Client ID#
1107985-35	1107985-35	1107985-35	1107985-35	1107985-34	1107985-34	1107985-34	1107985-34	1107985-33	1107985-33	1107985-33	1107985-33	1107985-32	1107985-32	1107985-32	1107985-32	Lab ID#
Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Sulfate	Nitrate	Nitrite	Chloride	Analyte
454	0.115	0.165	14.0	5500	0.115	0.165	41.7	146	0.115	0.165	6.40	1640	0.115	0.165	27.2	Result
3	_ ⊐	∪ m	Ħ	3	_ □	∪ m	3	æ	∪ m	□ m	3	3	⊂ m	⊂ m	3	N N
mg/L (mg/L (mg/L (mg/L (mg/L (Units										
0.200	0.115	0.165	0.200	0.200	0.115	0.165	0.200	0.200	0.115).165	0.200	0.200	0.115	0.165	0.200	<u>LOD</u>
2.5	0.6	0.8	2.5	2.5	0.6	0.8	2.5	2.5	0.6	0.8	2.5	2.5	0.6	0.8	2.5	<u> </u>
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_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	Matrix
300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	300.0	Method
KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	KMG	Analyst
12-May-11	12-May-11	12-May-11	12-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	03-May-11	Analyzed
21-Apr-11	21-Apr-1	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11	21-Apr-11		19-Apr-11			19-Apr-11	19-Apr-11	19-Apr-11	19-Apr-11	Collected

Phone: 330-253-8211

JPOP41S-110421

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit Env	rironmental Tech	Contract:	Jackpile	
Matrix: (soil/water) _	water	_ Lab Sample ID: _	1107985-01	
Date Received:	04/26/2011	_		
% Solids:				
		_		

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Q	М
7429-90-5	Aluminum	5.31			P
7440-36-0	Antimony	0.00237	U		Р
7440-38-2	Arsenic	0.00206	U		P
7440-39-3	Barium	0.008	J		P
7440-41-7	Beryllium	0.0395			Р
7440-43-9	Cadmium	0.0045	J		Р
7440-70-2	Calcium	744		D	Р
7440-47-3	Chromium	0.365			Р
7440-48-4	Cobalt	0.0788			Р
7440-50-8	Copper	0.00183	Ū		Р
7439-89-6	Iron	0.530	U	D	Р
7439-92-1	Lead	0.341			Р
7439-95-4	Magnesium	581		D	Р
7439-96-5	Manganese	2.84			Р
7440-02-0	Nickel	0.0086	J		Р
7440-09-7	Potassium	52.0			P
7782-49-2	Selenium	0.104			Р
7440-22-4	Silver	0.321			Р
7440-23-5	Sodium	561		D	Р
7440-28-0	Thallium	0.00529	Ū		Р
7440-62-2	Vanadium	0.00174			Р
7440-66-6	Zinc	0.0155	J		Р
7439-97-6	Mercury	0.00006	U		AV

Comment	S:
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_		D.	ue	to into	erterence:	s iron's	det	ection	limit	was	raised	and	reported	
	From	the	50	times	dilution	analyzed	on	05/12/	2011					
				-				*****						
					160									

Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-01 Run Time: 04/29/11 17:46:41

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0607	.5398	.0045	.3646	.0080	0105	.2509
SDev	.0042	.0020	.0002	.0028	.0003	.0001	.0025
%RSD	6.859	.3688	3.814	.7674	3.351	1.280	.9869
#1	0637	.5413	.0047	.3627	.0078	0106	.2491
#2	0578	.5384	.0044	.3666		0104	.2526
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0042	1493	.0155	.0086	0576	.0395	2113
SDev	.0029	.0006	.0003	.0008	.0019	.0001	.0097
%RSD	70.16	.4247	1.750	9.621	3.276	.2579	4.587
#1	0021	1498	.0157	.0092	0589	.0395	2044
#2	0062	1489	.0154		0562	.0396	2181
Elem Units Avge SDev %RSD	Sb206A ppm L-1.000 .007 .7261	S_1820 ppm H1077. 2. .1538	S_2203 ppm .3407 .0001 .0192	S_1960 ppm .1044 .0036 3.427	S_2068 ppm 7575 .0082 1.086	Ca3179 ppm H680.9 .8	Mg2790 ppm H491.6 .8 .1651
#1	L-1.000	H1076.	.3406	.1070	7517	H680.4	H491.1
#2	L-1.000	H1078.	.3407		7633	H681.5	H492.2
Elem Units Avge SDev %RSD	Mn2576 ppm 2.835 .003 .1096	Co2286 ppm .0788 .0009 1.099	Fe2714 ppm L-1.000 2 .031 7 .8058	Al3082 ppm 5.312 .050 .9395	V_2924 ppm 1734 .0028 1.604	Mo2020 ppm .0219 .0004 1.907	Ti3349 ppm .0739 .0001 .1211
#1	2.833	.0782	L-1.000	5.277	1754	.0222	.0739
#2	2.838	.0794	L-1.000	5.348	1715	.0216	.0740
Elem Units Avge SDev %RSD	Ag3280 ppm .3212 .0027 .8570	B_2496 ppm 1119 .0039 3.527	K_7664 ppm 51.97 .15 .2905	Na3302 ppm H683.7 .2	Si2881 ppm 5.389 .024 .4486	Sn1899 ppm 0428 .0017 3.909	P_1782 ppm .1597 .0044 2.765
#1	.3192	1147	52.08	H683.6	5.372	0440	.1628
#2		1091	51.86	H683.8	5.406	0416	.1566

Method: ZZJEM1_A Sample Name: 7985-01 x10 Operator: VVK

Run Time: 05/11/11 11:45:41

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Mode: CONC Corr. Factor: 1

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	1610	.1216	.0014	.1218	.0021	0116	.0337
SDev	.0181	.0034	.0004	.0008	.0001	.0022	.0042
%RSD	11.25	2.756	30.10	.6530	3.449	18.90	12.51
#1 #2	1482 1738	.1192 .1240	.0016	.1224	.0021	0131 0100	.0366 .0307
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0105	0171	.0377	.0494	0519	.0133	0064
SDev	.0123	.0007	.0002	.0018	.0143	.0000	.0002
%RSD	116.7	4.365	.4327	3.725	27.65	.0791	3.506
#1	0018	0177	.0376	.0507	0417	.0133	0063
#2	0192	0166	.0379	.0481	0620	.0133	0066
Elem Units Avge SDev %RSD	Sb206A ppm 0673 .0001 .1246	S_1820 ppm 60.27 .33 .5518	S_2203 ppm .0281 .0037 13.33	S_1960 ppm .0302 .0071 23.49	S_2068 ppm 0460 .0000	Ca3179 ppm 74.36 .55 .7446	Mg2790 ppm 58.05 .25 .4281
#1	0672	60.50	.0307	.0352	0459	74.75	58.23
#2	0674	60.03		.0252	0460	73.97	57.88
Elem Units Avge SDev %RSD	Mn2576 ppm .3547 .0025 .7160	Co2286 ppm .0136 .0004 3.202	Fe2714 ppm L-1.000 .058 2.770	Al3082 ppm 2.106 .003 .1584	V_2924 ppm 0303 .0009 3.011	Mo2020 ppm .0030 .0030 101.0	Ti3349 ppm .0156 .0002
#1 #2	.3565 .3529	.0133 .0139	L-1.000 L-1.000	2.108 2.103	0310 0297	.0051	.0155 .0157
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0424	0330	6.499	56.08	.6487	0021	.2688
SDev	.0001	.0005	.026	.14	.0004	.0002	.0004
%RSD	.3523	1.555	.4055	.2460	.0641	11.56	.1591
#1	.0423	0334	6.517	56.17	.6485	0020	.2691
#2	.0425	0326	6.480	55.98	.6490	0023	.2685

Method: ZZJEM1_A Sample Name: 7985-01 x50 Operator: VVK Run Time: 05/12/11 17:48:11

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0643	.0249	0010	.0279	0114	0000	.0146
SDev	.0124	.0041	.0004	.0002	.0001	.0080	.0037
%RSD	19.22	16.38	35.99	.7878	1.268	18290.	25.52
#1	0730	.0220	0008	.0278	0116	.0056	.0120
#2	0555	.0278	0013	.0281	0113	0057	.0173
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0099	0080	.0270	.0076	.0166	.0021	0110
SDev	.0049	.0006	.0062	.0016	.0018	.0001	.0012
%RSD	48.99	7.101	22.84	21.25	10.88	5.267	10.52
#1 #2	0134 0065	0084 0076	.0226 .0313	.0065	.0179 .0153	.0021	0102 0119
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0201	14.73	0043	.0248	0162	22.60	12.98
SDev	.0013	.05	.0069	.0045	.0004	.27	.06
%RSD	6.597	.3687	157.7	18.31	2.536	1.188	.4393
#1	0211	14.69	0092	.0216	0164	22.41	12.94
#2	0192	14.77	.0005		0159	22.79	13.02
Elem Units Avge SDev %RSD	Mn2576 ppm .0932 .0010 1.127	Co2286 ppm .0014 .0006 39.31	Fe2714 ppm 5164 .0305 5.913	Al3082 ppm .4588 .0143 3.110	V_2924 ppm 0096 .0004 4.532	Mo2020 ppm 0020 .0034 169.7	Ti3349 ppm .0018 .0003 17.67
#1 #2	.0925	.0010	5380 4948	.4487 .4688	0099 0093	.0004	.0016
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0086	.0002	1.000	6.164	.1583	0002	.0236
SDev	.0001	.0023	.064	.010	.0060	.0003	.0021
%RSD	1.740	945.9	6.436	.1645	3.791	134.9	8.706
#1 #2	.0087	0014 .0019	.9548 1.046	6.157 6.171	.1541 .1626	0004 0000	.0221

MD-SW-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit Env	rironmental Tech	Contract:	Jackpile	
Matrix: (soil/water) _	water	_ Lab Sample ID: _	1107985-02	
Date Received:	04/26/2011	- -		
% Solids:		_		

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Q	M
7429-90-5	Aluminum	0.0129	J		P
7440-36-0	Antimony	0.0079	J		Р
7440-38-2	Arsenic	0.0028	J		Р
7440-39-3	Barium	0.0722			Р
7440-41-7	Beryllium	0.0008	J	***************************************	P
7440-43-9	Cadmium	0.00082	U		Р
7440-70-2	Calcium	270		D	P
7440-47-3	Chromium	0.00109	U		Р
7440-48-4	Cobalt	0.0018	J		P
7440-50-8	Copper	0.00183	U		Р
7439-89-6	Iron	0.0106	Ū		P
7439-92-1	Lead	0.0026	J		Р
7439-95-4	Magnesium	198		D	P
7439-96-5	Manganese	0.151			Р
7440-02-0	Nickel	0.0036	J		P
7440-09-7	Potassium	26.8			Р
7782-49-2	Selenium	0.0322			Р
7440-22-4	Silver	0.0019	J		Р
7440-23-5	Sodium	1290		D	P
7440-28-0	Thallium	0.0062	J		Р
7440-62-2	Vanadium	0.00174	U		P
7440-66-6	Zinc	0.00053	U		Р
7439-97-6	Mercury	0.00006	U		AV
					

Comments:							
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Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-02 Run Time: 04/29/11 17:50:16

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0076 .0096 126.2	Pb220A ppm 0003 .0012 355.3	Cd2265 ppm .0008 .0002 23.31	Cr2677 ppm .0010 .0001 10.24	Ba4934 ppm .0722 .0000	As1890 ppm .0028 .0042 149.6	Se1960 ppm .0047 .0039 83.31
#1 #2	.0008	0012 .0005	.0010	.0010 .0009	.0722 .0722	.0058 0002	.0019
Elem Units Avge SDev %RSD	Se196A ppm .0001 .0039 7631.	Cu3247 ppm .0004 .0005 122.6	Zn2138 ppm 0094 .0000	Ni2316 ppm .0036 .0014 39.36	Tl1908 ppm .0062 .0026 42.04	Be3130 ppm .0008 .0001 16.77	Sb2068 ppm .0024 .0044 183.6
#1 #2	.0028 0027	.0007	0094 0094	.0046	.0043	.0008	.0055
Elem Units Avge SDev %RSD	Sb206A ppm .0090 .0031 34.28	S_1820 ppm H742.3 2.0 .2712	S_2203 ppm .0026 .0040 151.5	S_1960 ppm .0322 .0013 3.957	S_2068 ppm .0079 .0037 46.83	Ca3179 ppm H262.1 .3	Mg2790 ppm H174.8 \ .2 .1411
#1 #2	.0112	H740.9 H743.7	0002 .0055	.0331	.0105	H261.8 H262.3	H174.6 H174.9
Elem Units Avge SDev %RSD	Mn2576 ppm .1509 .0001 .0833	Co2286 ppm .0018 .0004 24.34	Fe2714 ppm 0146 .0001 .6888	Al3082 ppm .0129 .0104 80.36	V_2924 ppm 0606 .0003 .4507	Mo2020 ppm .0062 .0009 14.89	Ti3349 ppm0015 .0002 13.84
#1 #2	.1508 .1510	.0022 .0015	0145 0147	.0203	0604 0608	.0069	0014 0017
Elem Units Avge SDev %RSD	Ag3280 ppm .0019 .0012 64.97	B_2496 ppm .7973 .0029 .3677	K_7664 ppm 26.77 .04 .1334	Na3302 ppm H1819. 2.	Si2881 ppm 3.213 .002 .0773	Sn1899 ppm .0050 .0001 3.033	P_1782 ppm .0737 .0059 8.034
#1 #2	.0028	.7952 .7994	26.75 26.80	H1818. H1821.	3.215 3.211	.0049	.0779 .0695

Method: ZZJEM1_A Sample Name: 7985-02 x10 Operator: VVK

Run Time: 05/11/11 11:49:15

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0020	.0054	.0007	.0012	.0088	0049	0047
SDev	.0022	.0022	.0004	.0002	.0001	.0053	.0079
%RSD	107.2	40.78	60.28	18.89	.8270	108.8	168.1
#1 #2	.0036 .0005	.0069	.0004	.0010	.0088	0086 0011	0103 .0009
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908 ppm0332 .0044 13.31	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm		ppm	ppm
Avge	.0080	.0134	.0215	.0295		.0029	.0111
SDev	.0021	.0003	.0001	.0016		.0000	.0040
%RSD	26.08	2.605	.5324	5.432		1.445	35.86
#1	.0095	.0136	.0216	.0306	0300	.0029	.0083
#2	.0066	.0132	.0215	.0283	0363	.0029	.0139
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0023	37.61	.0045	.0293	.0061	26.95	19.75
SDev	.0040	.03	.0022	.0012	.0014	.11	.04
%RSD	178.6	.0905	49.16	3.949	22.25	.3931	.2012
#1 #2	.0051 0006	37.64 37.59	.0061	.0301 .0285	.0070	27.02 26.87	19.77 19.72
Elem	Mn2576	Co2286 ppm0002 .0003 139.6	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm		ppm	ppm	ppm	ppm	ppm
Avge	.0218		0096	.4426	0045	.0031	.0009
SDev	.0001		.0117	.0178	.0005	.0021	.0005
%RSD	.5512		122.6	4.024	10.27	69.32	53.88
#1 #2	.0219 .0217	0000 0004	0179 0013	.4300	0049 0042	.0046 .0016	.0013
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0003	.0492	3.751	H128.8	.4392	.0008	.1595
SDev	.0001	.0008	.006	.3	.0035	.0001	.0095
%RSD	31.16	1.544	.1729	.2543	.7895	9.111	5.956
#1	.0002	.0497	3.747	H129.1	.4368	.0007	.1528
#2		.0486	3.756	H128.6	.4417	.0009	.1662

MD-SW-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit Er	nvironmental Tech	_ Contract:	Jackpile	
Matrix: (soil/water)	water	_ Lab Sample ID: _	1107985-03	
Date Received:	04/26/2011			
% Solids:				

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Q	М
7429-90-5	Aluminum	0.0240	J		P
7440-36-0	Antimony	0.0075	J		Р
7440-38-2	Arsenic	0.00206	Ū		P
7440-39-3	Barium	0.0180	J		P
7440-41-7	Beryllium	0.00072	U		P
7440-43-9	Cadmium	0.00082	Ū		P
7440-70-2	Calcium	28.7		D	Р
7440-47-3	Chromium	0.00109	U		Р
7440-48-4	Cobalt	0.0022	J		Р
7440-50-8	Copper	0.00183	Ū		Р
7439-89-6	Iron	0.0106	Ū		Р
7439-92-1	Lead	0.00210	U		P
7439-95-4	Magnesium	85.54			Р
7439-96-5	Manganese	0.00043	U		P
7440-02-0	Nickel	0.0025	J		Р
7440-09-7	Potassium	7.031		N	P
7782-49-2	Selenium	0.0309			Р
7440-22-4	Silver	0.0014	J	N	P
7440-23-5	Sodium	867		D	Р
7440-28-0	Thallium	0.0212			P
7440-62-2	Vanadium	0.00174	U		P
7440-66-6	Zinc	0.00053	Ū		P
7439-97-6	Mercury	0.00006	Ū		AV

Comn	nents	3:										
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Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-03 Run Time: 04/29/11 17:53:50

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0009 .0038 421.3	Pb220A ppm 0008 .0003 37.46	Cd2265 ppm .0005 .0002 38.68	Cr2677 ppm .0004 .0000 4.024	Ba4934 ppm .0180 .0000	As1890 ppm 0010 .0047 455.7	Se1960 ppm .0018 .0064 359.0
#1 #2	0018 .0036	0006 0010	.0006	.0004	.0180	.0023 0044	0027 .0063
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm						
Avge	.0022	0018	0036	.0025	.0212	.0003	.0044
SDev	.0063	.0010	.0001	.0004	.0040	.0001	.0026
%RSD	290.1	55.00	1.610	16.01	18.93	15.74	59.65
#1	.0066	0025	0037	.0022	.0240	.0004	.0062
#2	0023	0011	0036	.0028	.0183		.0025
Elem Units Avge SDev %RSD	Sb206A ppm .0076 .0020 25.84	S_1820 ppm H401.1 .5 .1331	S_2203 ppm .0001 .0011 872.2	S_1960 ppm .0309 .0044 14.34	S_2068 ppm .0075 .0022 29.01	Ca3179 ppm 24.53 .00 .0199	Mg2790 ppm 85.54 .16
#1 #2	.0090 .0062	H400.7 H401.4	0006 .0009	.0341	.0091	24.53 24.54	85.43 85.65
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm						
Avge	0173	.0022	0308	.0240	0444	.0069	0003
SDev	.0001	.0005	.0158	.0010	.0003	.0002	.0001
%RSD	.5256	23.55	51.23	4.288	.7472	2.794	20.95
#1	0172	.0019	0419	.0233	0442	.0071	0003
#2	0174		0196	.0247	0446	.0068	0002
Elem Units Avge SDev %RSD	Ag3280 ppm .0014 .0004 31.00	B_2496 ppm .4770 .0016 .3433	K_7664 ppm 7.031 .068 .9657	Na3302 ppm H1284. 2.	Si2881 ppm 3.574 .003 .0707	Sn1899 ppm .0080 .0026 32.18	P_1782 ppm .0692 .0051 7.389
#1	.0011	.4782	6.983	H1283.	3.573	.0061	.0656
#2		.4759	7.079	H1286.	3.576	.0098	.0728

Method: ZZJEM1_A Sample Name: 7985-03 x10 Operator: VVK

Run Time: 05/11/11 11:52:49

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0068	.0052	0000	.0013	.0026	0045	0024
SDev	.0003	.0045	.0000	.0003	.0001	.0048	.0102
%RSD	3.746	86.28	148.1	26.52	2.773	106.4	424.6
#1	0066	.0020	0001	.0015	.0026	0011	.0048
#2	0069	.0084	.0000		.0027	0080	0096
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0042	.0068	.0148	.0179	0082	.0027	.0050
SDev	.0008	.0003	.0001	.0011	.0055	.0002	.0014
%RSD	19.09	3.683	.3867	5.948	67.34	5.813	28.89
#1 #2	.0048 .0037	.0069	.0148	.0187 .0172	0043 0121	.0028	.0039
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0054	19.60	.0016	.0309	.0062	2.870	9.392
SDev	.0023	.02	.0029	.0015	.0010	.019	.004
%RSD	41.96	.1086	180.5	4.869	16.54	.6592	.0393
#1 #2	.0070 .0038	19.61 19.58	0004 .0037	.0320	.0070	2.856 2.883	9.390 9.395
Elem Units Avge SDev %RSD	Mn2576 ppm .0009 .0001 6.409	Co2286 ppm .0008 .0005 66.01	Fe2714 ppm 0196 .0122 62.20	Al3082 ppm .4398 .0007 .1541	V_2924 ppm 0028 .0004 15.98	Mo2020 ppm 0002 .0005 212.1	Ti3349 ppm .0003 .0000
#1 #2	.0009	.0004	0282 0110	.4393	0025 0031	.0001 0006	.0003
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0006	.0062	1.478	86.73	.4906	0002	.0614
SDev	.0003	.0006	.007	.29	.0046	.0010	.0078
%RSD	62.32	9.764	.5003	.3380	.9330	475.3	12.69
#1 #2	.0003	.0058 .0066	1.483 1.473	86.93 86.52	.4874 .4939	0009 .0005	.0559

MW-2-110421

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name:Summit Environmental Tech	Contract:	Jackpile
Matrix: (soil/water) water	Lab Sample ID:	1107985-04
Date Received: 04/26/2011		2207903 01
% Solids:		
Concentration Units (mg/L or mg/kg dry weight	:): mq/L	

CAS No.	Analyte	Concentration	С	Q	М
7429-90-5	Aluminum	0.0163	J		Р
7440-36-0	Antimony	0.0061	J		Р
7440-38-2	Arsenic	0.0033	J		P
7440-39-3	Barium	0.0207	J		Р
7440-41-7	Beryllium	0.00072	U		P
7440-43-9	Cadmium	0.00082	U		P
7440-70-2	Calcium	247		D	P
7440-47-3	Chromium	0.00109	U		P
7440-48-4	Cobalt	0.0006	J		P
7440-50-8	Copper	0.00183	U		P
7439-89-6	Iron	1.15			P
7439-92-1	Lead	0.0030	J		P
7439-95-4	Magnesium	279		D	P
7439-96-5	Manganese	0.275			Р
7440-02-0	Nickel	0.0048	J		P
7440-09-7	Potassium	18.7			P
7782-49-2	Selenium	0.0317			P
7440-22-4	Silver	0.0020	J		P
7440-23-5	Sodium	648		D	P
7440-28-0	Thallium	0.0106	J		Р
7440-62-2	Vanadium	0.00174	U		P
7440-66-6	Zinc	0.0017	J		P
7439-97-6	Mercury	0.00006	U		AV

Com	ment:	s:							
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Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-04 Run Time: 04/29/11 18:04:31

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0076 .0011 14.02	Pb220A ppm .0004 .0004 117.6	Cd2265 ppm .0004 .0002 43.44	Cr2677 ppm .0003 .0003 110.0	Ba4934 ppm .0207 .0001 .2605	As1890 ppm .0033 .0025 76.72	Se1960 ppm .0034 .0052 154.0
#1 #2	.0084	.0007	.0006	.0001 .0006	.0207	.0015 .0050	0003 .0071
Elem Units Avge SDev %RSD	Se196A ppm .0027 .0089 336.0	Cu3247 ppm 0025 .0008 30.50	Zn2138 ppm .0017 .0003 16.44	Ni2316 ppm .0048 .0011 22.80	Tl1908 ppm .0106 .0233 220.3	Be3130 ppm .0006 .0000 3.908	Sb2068 ppm .0025 .0015 59.73
#1 #2	.0090 0037	0020 0031	.0019 .0015	.0056 .0041	.0270 0059	.0006	.0036 .0015
Elem Units Avge SDev %RSD	Sb206A ppm .0063 .0040 62.94	S_1820 ppm H701.3 1.4 .1928	S_2203 ppm .0030 .0006 20.71	S_1960 ppm .0317 .0042 13.33	S_2068 ppm .0061 .0022 35.54	Ca3179 ppm H247.0 .6	Mg2790 ppm H256.0 .7 .2738
#1 #2	.0035	H700.3 H702.2	.0034	.0347 .0287	.0045	H246.6 H247.4	H255.5 H256.5
Elem Units Avge SDev %RSD	Mn2576 ppm .2747 .0005 .1947	Co2286 ppm .0006 .0004 66.21	Fe2714 ppm 1.154 .005 .4228	Al3082 ppm .0163 .0005 2.961	V_2924 ppm 0325 .0006 1.901	Mo2020 ppm .0086 .0005 6.232	Ti3349 ppm 0015 .0001 5.851
#1 #2	.2743 .2751	.0009	1.158 1.151	.0167 .0160	0321 0330	.0082	0015 0016
Elem Units Avge SDev %RSD	Ag3280 ppm .0020 .0018 89.46	B_2496 ppm .2158 .0007 .3406	K_7664 ppm 18.73 .11 .5617	Na3302 ppm H942.5 4.0 .4249	Si2881 ppm 9.032 .025 .2742	Sn1899 ppm .0042 .0013 30.39	P_1782 ppm .0774 .0008 1.068
#1 #2	.0007	.2163 .2152	18.65 18.80	H939.6 H945.3	9.014 9.049	.0051	.0768 .0780

Method: ZZJEM1_A Sample Name: 7985-04 x10 Operator: VVK Run Time: 05/11/11 11:56:23

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0033 .0001	Pb220A ppm .0041 .0021 52.46	Cd2265 ppm .0006 .0006 95.81	Cr2677 ppm .0011 .0001 10.13	Ba4934 ppm .0028 .0001 5.238	As1890 ppm 0040 .0031 75.64	Se1960 ppm .0065 .0139 213.8
#1 #2	.0033	.0056 .0026	.0002	.0010 .0012	.0029	0019 0062	0033 .0164
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908 ppm .0004 .0386 9899.	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm		ppm	ppm
Avge	.0034	.0055	.0058	.0080		.0029	.0054
SDev	.0010	.0001	.0001	.0015		.0002	.0028
%RSD	28.57	1.825	2.047	19.35		5.464	51.81
#1 #2	.0041	.0054	.0057 .0059	.0069 .0091	.0277 0269	.0028	.0034
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm						
Avge	0004	35.34	.0041	.0352	.0026	24.70	27.87
SDev	.0023	.12	.0014	.0040	.0025	.17	.08
%RSD	660.0	.3330	34.23	11.28	97.65	.6790	.2751
#1 #2	0020 .0013	35.25 35.42	.0051	.0324	.0008	24.58 24.82	27.82 27.93
Elem	Mn2576	Co2286	Fe2714	A13082	V_2924	Mo2020	Ti3349
Units	ppm						
Avge	.0342	0004	.0585	.4500	0024	.0006	0002
SDev	.0003	.0008	.0134	.0007	.0000	.0010	.0004
%RSD	.9277	219.7	22.94	.1467	.0551	169.7	169.7
#1	.0339	0009	.0680	.4505	0024	.0013	.0000
#2		.0002	.0490	.4496	0024	0001	0005
Elem Units Avge SDev %RSD	Ag3280 ppm .0000 .0003 1116.	B_2496 ppm 0266 .0014 5.256	K_7664 ppm 1.776 .009	Na3302 ppm 64.84 .04 .0591	Si2881 ppm 1.134 .003 .2568	Sn1899 ppm .0003 .0010 355.4	P_1782 ppm .0426 .0068 15.95
#1	0002	0256	1.769	64.81	1.132	0004	.0378
#2	.0003	0276	1.782	64.87	1.137	.0010	.0474

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit En	vironmental Tech	Contract:	Jackpile	
Matrix: (soil/water)	water	_ Lab Sample ID:	1107985-05	
Date Received:	04/26/2011	_		
% Solids:	4	_		

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Õ	M
7429-90-5	Aluminum	0.0164	J		Р
7440-36-0	Antimony	0.0057	J		P
7440-38-2	Arsenic	0.00206	U		P
7440-39-3	Barium	0.0221	J		P
7440-41-7	Beryllium	0.00072	Ū		Р
7440-43-9	Cadmium	0.00082	Ū		Р
7440-70-2	Calcium	267		D	Р
7440-47-3	Chromium	0.0019	U		P
7440-48-4	Cobalt	0.0006	J		Р
7440-50-8	Copper	0.00183	Ū		Р
7439-89-6	Iron	1.27			P
7439-92-1	Lead	0.00210	U		Р
7439-95-4	Magnesium	297		D	Р
7439-96-5	Manganese	0.294			Р
7440-02-0	Nickel	0.0052	J		Р
7440-09-7	Potassium	20.32			Р
7782-49-2	Selenium	0.0327		1	Р
7440-22-4	Silver	0.00039	U		Р
7440-23-5	Sodium	694		D	Р
7440-28-0	Thallium	0.007	J		Р
7440-62-2	Vanadium	0.00174	U		Р
7440-66-6	Zinc	0.00053	U		Р
7439-97-6	Mercury	0.00006	U		AV

Comments	:						
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Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-05 Run Time: 04/29/11 18:08:05

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0023 .0032 139.9	Pb220A ppm 0012 .0001 12.02	Cd2265 ppm .0003 .0001 43.37	Cr2677 ppm .0000 .0007 3052.	Ba4934 ppm .0221 .0001 .2434	As1890 ppm .0017 .0004 22.23	Se1960 ppm .0061 .0015 24.88
#1 #2	.0000	0011 0014	.0004	.0005 0004	.0221	.0019	.0050
Elem Units Avge SDev %RSD	Se196A ppm .0026 .0020 77.61	Cu3247 ppm 0025 .0010 40.61	Zn2138 ppm0005 .0001 10.95	Ni2316 ppm .0052 .0010 20.29	Tl1908 ppm .0077 .0111 143.5	Be3130 ppm .0004 .0001 16.65	Sb2068 ppm .0038 .0013 35.50
#1 #2	.0012	0018 0032	0004 0005	.0059	.0156 0001	.0004	.0028
Elem Units Avge SDev %RSD	Sb206A ppm .0052 .0021 39.56	S_1820 ppm H739.4 1.5 .2042	S_2203 ppm .0001 .0008 598.1	S_1960 ppm .0327 .0018 5.545	S_2068 ppm .0057 .0009 16.30	Ca3179 ppm H262.9 .6	Mg2790 ppm H274.1 .7 .2401
#1 #2	.0067	H738.3 H740.5	0004 .0007	.0315	.0064	H262.5 H263.4	H273.6 H274.6
Elem Units Avge SDev %RSD	Mn2576 ppm .2940 .0008 .2880	Co2286 ppm .0006 .0000 6.270	Fe2714 ppm 1.267 .018 1.449	Al3082 ppm .0164 .0000 .1103	V_2924 ppm 0350 .0005 1.338	Mo2020 ppm .0077 .0008 10.42	Ti3349 ppm 0020 .0004 20.73
#1 #2	.2934 .2945	.0006	1.280 1.255	.0164 .0164	0346 0353	.0083	0017 0023
Elem Units Avge SDev %RSD	Ag3280 ppm 0001 .0010 677.2	B_2496 ppm .2224 .0011 .4926	K_7664 ppm 20.32 .03 .1424	Na3302 ppm H1012. 3. .2708	Si2881 ppm 9.611 .028 .2879	Sn1899 ppm .0024 .0021 86.67	P_1782 ppm .0852 .0025 2.911
#1 #2	.0005 0008	.2216 .2232	20.30 20.34	H1011. H1014.	9.592 9.631	.0038	.0834

Method: ZZJEM1_A Sample Name: 7985-05 x10 Operator: VVK Run Time: 05/11/11 11:59:57 Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0032 .0014 43.13	Pb220A ppm .0058 .0030 52.53	Cd2265 ppm .0000 .0004 1683.	Cr2677 ppm .0009 .0004 51.61	Ba4934 ppm .0029 .0001 2.481	As1890 ppm 0100 .0040 39.65	Se1960 ppm 0002 .0047 1901.
#1 #2	.0041	.0079 .0036	.0003	.0012	.0030	0072 0128	.0031 0036
Elem Units Avge SDev %RSD	Se196A ppm 0013 .0021 159.3	Cu3247 ppm .0063 .0001 .7857	Zn2138 ppm .0075 .0000 .0643	Ni2316 ppm .0106 .0002 2.245	Tl1908 ppm 0055 .0425 777.8	Be3130 ppm .0029 .0000 1.219	Sb2068 ppm .0043 .0032 74.25
#1 #2	0027 .0002	.0063	.0075	.0104	.0246 0355	.0029	.0065
Elem Units Avge SDev %RSD	Sb206A ppm .0011 .0033 290.3	S_1820 ppm 37.11 .05 .1473	S_2203 ppm .0052 .0025 48.40	S_1960 ppm .0264 .0050 19.00	S_2068 ppm .0031 .0010 33.39	Ca3179 ppm 26.65 .23 .8519	Mg2790 ppm 29.66 .02
#1 #2	0012 .0034	37.15 37.07	.0069	.0299 .0228	.0024	26.49 26.81	29.65 29.67
Elem Units Avge SDev %RSD	Mn2576 ppm .0365 .0001 .1525	Co2286 ppm .0002 .0004 216.8	Fe2714 ppm .0615 .0019 3.142	Al3082 ppm .4673 .0000	V_2924 ppm 0022 .0009 42.00	Mo2020 ppm .0015 .0007 47.14	Ti3349 ppm .0002 .0003 176.7
#1 #2	.0365 .0365	.0005 0001	.0601 .0628	.4673 .4673	0015 0028	.0019 .0010	.0004
Elem Units Avge SDev %RSD	Ag3280 ppm .0003 .0002 62.30	B_2496 ppm 0250 .0006 2.368	K_7664 ppm 2.112 .006 .2843	Na3302 ppm 69.40 .37 .5355	Si2881 ppm 1.200 .001 .0924	Sn1899 ppm .0008 .0003 32.94	P_1782 ppm .0709 .0012 1.676
#1 #2	.0002 .0005	0246 0255	2.108 2.117	69.14 69.67	1.200	.0006	.0700 .0717

MW-3-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name:	Summit Envir	onmental Te	ech Contr	act:		Jackpil	le
Matrix: (soi	l/water)	water	Lab S	ample	ID: _	110798	35-06
Date Received	d:	04/26/2011					
% Solids:							
Concentration	n Units (mg/	L or mg/kg	dry weight): <u>m</u>	g/L	_		
	CAS No.	Analyte	Concentration	С		Q	M
			<u> </u>	 			

CAS No.	Analyte	Concentration	С	Q	M
7429-90-5	Aluminum	0.0152	J		Р
7440-36-0	Antimony	0.0024	J		P
7440-38-2	Arsenic	0.00206	U		P
7440-39-3	Barium	0.0281	J		P
7440-41-7	Beryllium	0.00072	Ū		Р
7440-43-9	Cadmium	0.00082	Ū		Р
7440-70-2	Calcium	192		D	Р
7440-47-3	Chromium	0.00109	U		Р
7440-48-4	Cobalt	0.00051	Ū		Р
7440-50-8	Copper	0.00183	U		Р
7439-89-6	Iron	0.0106	U		Р
7439-92-1	Lead	0.00210	Ū		Р
7439-95-4	Magnesium	57.5			Р
7439-96-5	Manganese	0.00043	U		P
7440-02-0	Nickel	0.0016	U		P
7440-09-7	Potassium	5.92		N	P
7782-49-2	Selenium	0.0337			P
7440-22-4	Silver	0.00039	U	N	P
7440-23-5	Sodium	81.0		D	P
7440-28-0	Thallium	0.0059	U		P
7440-62-2	Vanadium	0.00174	U		P
7440-66-6	Zinc	0.0066	J		Р
7439-97-6	Mercury	0.00006	U		AV

Comme	ents:								
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Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-06 Run Time: 04/29/11 18:11:39

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0055 .0086 156.6	Pb220A ppm 0043 .0031 72.19	Cd2265 ppm .0000 .0000 59.17	Cr2677 ppm .0004 .0001 18.85	Ba4934 ppm .0281 .0000	As1890 ppm 0013 .0000 .9999	Se1960 ppm .0163 .0205 126.1
#1 #2	0006 .0116	0065 0021	.0000	.0004	.0281 .0281	0014 0013	.0307
Elem Units Avge SDev %RSD	Se196A ppm .0042 .0025 59.26	Cu3247 ppm 0026 .0001 4.119	Zn2138 ppm .0066 .0001 1.322	Ni2316 ppm .0007 .0007 110.8	Tl1908 ppm .0019 .0034 173.7	Be3130 ppm .0003 .0001 37.05	Sb2068 ppm 0025 .0056 221.4
#1 #2	.0025	0025 0026	.0065 .0067	.0001	.0043 0004	.0003	0065 .0014
Elem Units Avge SDev %RSD	Sb206A ppm .0035 .0079 226.7	S_1820 ppm H165.8 .4 .2428	S_2203 ppm 0007 .0049 700.8	S_1960 ppm .0337 .0027 8.129	S_2068 ppm .0024 .0033 136.4	Ca3179 ppm H191.8 .4 .2215	Mg2790 ppm 57.48 .17 .2917
#1 #2	.0090 0021	H165.5 H166.1	0042 .0028	.0357 .0318	.0047	H191.5 H192.1	57.36 57.60
Elem Units Avge SDev %RSD	Mn2576 ppm 0013 .0001 5.806	Co2286 ppm0000 .0003 648.5	Fe2714 ppm0402 .0027 6.657	Al3082 ppm .0152 .0076 50.06	V_2924 ppm 0040 .0003 6.926	Mo2020 ppm .0009 .0001 8.222	Ti3349 ppm0018 .0002 9.371
#1 #2	0012 0013	.0002 0003	0421 0383	.0206 .0098	0042 0038	.0009 .0010	0019 0016
Elem Units Avge SDev %RSD	Ag3280 ppm .0003 .0002 75.08	B_2496 ppm .0447 .0013 2.981	K_7664 ppm 5.915 .075 1.263	Na3302 ppm H103.9 .3	Si2881 ppm 8.209 .022 .2678	Sn1899 ppm .0100 .0003 3.444	P_1782 ppm .0868 .0020 2.274
#1 #2	.0001	.0437	5.968 5.862	H104.1 H103.7	8.193 8.224	.0098	.0854

Method: ZZJEM1_A Sample Name: 7985-06 x10 Operator: VVK Run Time: 05/11/11 12:03:31

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0083 .0010 12.58	Pb220A ppm .0068 .0008 12.24	Cd2265 ppm .0004 .0003 89.49	Cr2677 ppm .0002 .0003 212.3	Ba4934 ppm .0037 .0000	As1890 ppm 0138 .0005 3.848	Se1960 ppm .0060 .0115 189.7
#1 #2	.0076	.0074	.0006	0001 .0004	.0037	0142 0134	0021 .0142
Elem Units Avge SDev %RSD	Se196A ppm 0025 .0062 247.9	Cu3247 ppm .0047 .0004 8.507	Zn2138 ppm .0045 .0002 3.897	Ni2316 ppm .0066 .0002 3.580	Tl1908 ppm 0261 .0276 105.5	Be3130 ppm .0029 .0001 4.169	Sb2068 ppm .0059 .0020 33.43
#1 #2	0068 .0019	.0050	.0043	.0068	0066 0456	.0028	.0045
Elem Units Avge SDev %RSD	Sb206A ppm 0008 .0026 337.2	S_1820 ppm 8.147 .008 .0992	S_2203 ppm .0076 .0002 2.776	S_1960 ppm .0259 .0055 21.38	S_2068 ppm .0024 .0011 44.32	Ca3179 ppm 19.19 .04 .2103	Mg2790 ppm 6.406 .009 .1445
#1 #2	.0011 0026	8.153 8.141	.0077	.0220 .0298	.0032	19.22 19.16	6.399 6.412
Elem Units Avge SDev %RSD	Mn2576 ppm .0010 .0001 12.69	Co2286 ppm .0005 .0006 141.5	Fe2714 ppm 0188 .0027 14.24	Al3082 ppm .4701 .0158 3.369	V_2924 ppm .0004 .0002 62.09	Mo2020 ppm .0017 .0005 30.30	Ti3349 ppm0000 .0004 3183000.
#1 #2	.0011	.0009	0206 0169	.4589	.0002	.0021	.0003
Elem Units Avge SDev %RSD	Ag3280 ppm 0001 .0002 134.6	B_2496 ppm 0487 .0014 2.802	K_7664 ppm .7570 .0097 1.283	Na3302 ppm 8.103 .033	Si2881 ppm 1.059 .002 .1572	Sn1899 ppm .0006 .0005 84.81	P_1782 ppm .0264 .0045 16.93
#1 #2	0002 0000	0477 0497	.7502 .7639	8.126 8.080	1.060 1.058	.0010	.0232 .0295

MW-4-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit Env	rironmental Tech	Contract:	Jackpile	
Matrix: (soil/water) _	water	Lab Sample ID:	1107985-07	
Date Received:	04/26/2011	_		
% Solids:		_		

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Õ	М
7429-90-5	Aluminum	0.0117	J		Р
7440-36-0	Antimony	0.0207			P
7440-38-2	Arsenic	0.0041	J		Р
7440-39-3	Barium	0.0357	J		Р
7440-41-7	Beryllium	0.0008	J		Р
7440-43-9	Cadmium	0.00082	U		Р
7440-70-2	Calcium	203		D	Р
7440-47-3	Chromium	0.00109	U		P
7440-48-4	Cobalt	0.0011	J	200000000000000000000000000000000000000	Р
7440-50-8	Copper	0.00183	Ū		Р
7439-89-6	Iron	0.0106	U		Р
7439-92-1	Lead	0.0027	J		Р
7439-95-4	Magnesium	121		D	Р
7439-96-5	Manganese	0.0324			Р
7440-02-0	Nickel	0.0022	J		P
7440-09-7	Potassium	11.8			Р
7782-49-2	Selenium	0.0395			Р
7440-22-4	Silver	0.0025	J		Р
7440-23-5	Sodium	152		D	Р
7440-28-0	Thallium	0.0075	J		Р
7440-62-2	Vanadium	0.001741	Ū		Р
7440-66-6	Zinc	0.0103	J		P
7439-97-6	Mercury	0.00006	U		AV

Com	ments:					

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Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-07 Run Time: 04/29/11 18:29:30 Comment: 6010/200.7 CPI 4400-013 lot 10B101

Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0072 .0019 26.23	Pb220A ppm 0001 .0009 1226.	Cd2265 ppm .0007 .0001 11.73	Cr2677 ppm .0007 .0002 34.28	Ba4934 ppm .0357 .0001 .1506	As1890 ppm .0041 .0007 18.13	Se1960 ppm .0132 .0044 33.44
#1 #2	.0086	0007 .0006	.0007	.0005	.0358 .0357	.0046	.0101
Elem Units Avge SDev %RSD	Se196A ppm .0094 .0058 61.92	Cu3247 ppm .0002 .0003 113.1	Zn2138 ppm .0103 .0003 2.728	Ni2316 ppm .0022 .0009 41.64	Tl1908 ppm .0075 .0067 89.04	Be3130 ppm .0008 .0000 1.441	Sb2068 ppm .0174 .0079 45.21
#1 #2	.0135	.0004	.0105	.0029 .0016	.0122 .0028	.0008	.0230 .0119
Elem Units Avge SDev %RSD	Sb206A ppm .0208 .0053 25.69	S_1820 ppm H246.7 .1 .0265	S_2203 ppm .0027 .0000 1.265	S_1960 ppm .0395 .0048 12.14	S_2068 ppm .0207 .0062 29.94	Ca3179 ppm H202.6 .1 .0306	Mg2790 ppm H110.5 .0
#1 #2	.0246 .0170	H246.8 H246.7	.0028	.0429	.0250 .0163	H202.5 H202.6	H110.4 H110.5
Elem Units Avge SDev %RSD	Mn2576 ppm .0324 .0001 .1946	Co2286 ppm .0011 .0004 36.56	Fe2714 ppm 0496 .0121 24.38	Al3082 ppm .0117 .0020 17.35	V_2924 ppm 0063 .0001 2.074	Mo2020 ppm .0045 .0010 21.48	Ti3349 ppm0009 .0001 11.51
#1 #2	.0324	.0014 .0008	0581 0410	.0103 .0132	0062 0064	.0052	0008 0010
Elem Units Avge SDev %RSD	Ag3280 ppm .0025 .0007 27.08	B_2496 ppm .0849 .0018 2.089	K_7664 ppm 11.75 .08 .6862	Na3302 ppm H203.8 .9 .4229	Si2881 ppm 9.815 .002 .0206	Sn1899 ppm .0054 .0000 .4316	P_1782 ppm .0906 .0013 1.411
#1 #2	.0030	.0862 .0837	11.81 11.69	H204.4 H203.2	9.813 9.816	.0054	.0897 .0915

Method: ZZJEM1_A Sample Name: 7985-07 x10 Operator: VVK Run Time: 05/11/11 12:07:04

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm 0011 .0116 1078.	Pb220A ppm .0031 .0048 153.3	Cd2265 ppm .0004 .0000	Cr2677 ppm .0014 .0006 39.30	Ba4934 ppm .0044 .0000	As1890 ppm 0049 .0006 12.12	Se1960 ppm .0040 .0096 240.8
#1 #2	0093 .0071	0003 .0065	.0004	.0010	.0044	0045 0053	.0108 0028
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0052	.0055	.0070	.0081	0133	.0030	.0070
SDev	.0061	.0002	.0001	.0027	.0237	.0001	.0035
%RSD	117.9	3.650	1.745	33.53	178.9	1.538	50.74
#1 #2	.0096	.0056 .0053	.0069	.0062	-,0300 .0035	.0031	.0045 .0095
Elem Units Avge SDev %RSD	Sb206A ppm 0009 .0008 84.85	S_1820 ppm 12.03 .00	S_2203 ppm .0021 .0070 337.0	S_1960 ppm .0303 .0097 32.09	S_2068 ppm .0028 .0008 27.36	Ca3179 ppm 20.26 .05	Mg2790 ppm 12.07 .00
#1	0004	12.03	0029	.0372	.0023	20.23	12.07
#2	0014	12.03	.0070	.0234		20.29	12.07
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0051	.0001	0367	.4981	0002	.0013	.0004
SDev	.0000	.0007	.0099	.0092	.0007	.0020	.0003
%RSD	.1216	711.3	26.92	1.850	344.4	154.9	62.85
#1	.0051	0004	0297	.5046	.0003	.0027	.0006
#2	.0051	.0006	0437	.4916	0007	0001	
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0002	0454	1.364	15.22	1.234	.0006	.0404
SDev	.0003	.0002	.010	.29	.002	.0001	.0082
%RSD	152.2	.3898	.7118	1.901	.1462	19.06	20.35
#1	.0005	0456	1.371	15.43	1.232	.0006	.0462
#2	0000	0453	1.357	15.02	1.235	.0007	.0346

MW-6-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name:	Summit Env	ironmental Tech	Contract:	Jackpile	
Matrix: (soi	l/water) _	water	Lab Sample ID:	1107985-08	
Date Receive	d:	04/26/2011			
% Solids:					

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Q	М
7429-90-5	Aluminum	0.0161	J		P
7440-36-0	Antimony	0.0056	J		Р
7440-38-2	Arsenic	0.00206	Ū		Р
7440-39-3	Barium	0.0148	J		P
7440-41-7	Beryllium	0.00072	U		Р
7440-43-9	Cadmium	0.00082	U		P
7440-70-2	Calcium	278		D	Р
7440-47-3	Chromium	0.00109	U		P
7440-48-4	Cobalt	0.00051	U		Р
7440-50-8	Copper	0.00183	U		P
7439-89-6	Iron	0.0924	J		Р
7439-92-1	Lead	0.00210	Ū		Р
7439-95-4	Magnesium	205		D	Р
7439-96-5	Manganese	0.0193	J		Р
7440-02-0	Nickel	0.0020	J		Р
7440-09-7	Potassium	16.9			Р
7782-49-2	Selenium	0.0288			P
7440-22-4	Silver	0.0010	J		P
7440-23-5	Sodium	450		D	Р
7440-28-0	Thallium	0.0081	J		Р
7440-62-2	Vanadium	0.00174	U		Р
7440-66-6	Zinc	0.0037	J		Р
7439-97-6	Mercury	0.00006	U		AV

Comments:			
	No		

Method: ZZJEM1_A Sample Name: 7985-08 Run Time: 04/29/11 18:33:04

Elem Units Avge SDev %RSD	Pb2203 ppm .0046 .0020 42.80	Pb220A ppm .0002 .0020 1258.	Cd2265 ppm .0003 .0001 21.21	Cr2677 ppm .0004 .0005 142.0	Ba4934 ppm .0148 .0002 1.091	As1890 ppm .0009 .0001 6.011	Se1960 ppm .0013 .0019 146.2
#1 #2	.0060 .0032	.0016 0013	.0002	.0008	.0149 .0147	.0009	0000 .0026
Elem Units Avge SDev %RSD	Se196A ppm .0016 .0001 4.021	Cu3247 ppm 0016 .0002 13.26	Zn2138 ppm .0037 .0003 8.366	Ni2316 ppm .0020 .0010 47.41	Tl1908 ppm .0081 .0004 4.877	Be3130 ppm .0004 .0000 9.029	Sb2068 ppm .0030 .0024 79.64
#1 #2	.0016	0017 0014	.0039	.0014	.0083	.0004	.0047
Elem Units Avge SDev %RSD	Sb206A ppm .0054 .0001 .9686	S_1820 ppm H561.4 3.2 .5747	S_2203 ppm .0019 .0020 102.2	S_1960 ppm .0288 .0006 2.004	S_2068 ppm .0056 .0008	Ca3179 ppm H265.0 1.3 .5016	Mg2790 ppm H183.8 1.0 .5239
#1 #2	.0054	H563.7 H559.1	.0034	.0284	.0061 .0050	H266.0 H264.1	H184.5 H183.1
Elem Units Avge SDev %RSD	Mn2576 ppm .0193 .0001 .3845	Co2286 ppm .0002 .0002 86.88	Fe2714 ppm .0924 .0021 2.248	Al3082 ppm .0161 .0018 10.92	V_2924 ppm 0217 .0005 2.226	Mo2020 ppm .0052 .0011 21.06	Ti3349 ppm 0017 .0002 10.67
#1 #2	.0193 .0192	.0003	.0939	.0149 .0173	0221 0214	.0060	0016 0018
Elem Units Avge SDev %RSD	Ag3280 ppm .0010 .0003 30.34	B_2496 ppm .1714 .0026 1.502	K_7664 ppm 16.94 .05 .2975	Na3302 ppm H635.5 3.2 .5045	Si2881 ppm 8.596 .056 .6462	Sn1899 ppm .0037 .0012 33.21	P_1782 ppm .0773 .0004 .4862
#1 #2	.0008	.1732 .1696	16.97 16.90	Н637.8 Н633.3	8.636 8.557	.0045	.0776 .0770

Method: ZZJEM1_A Sample Name: 7985-08 x10 Operator: VVK Run Time: 05/11/11 12:10:38

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0065	.0008	.0002	.0009	.0021	0070	.0036
SDev	.0042	.0023	.0001	.0007	.0001	.0040	.0059
%RSD	65.08	294.6	45.37	77.17	3.449	56.57	163.9
#1 #2	.0094	.0024	.0001	.0004	.0021 .0022	0042 0098	0006 .0077
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0037	.0050	.0046	.0068	0258	.0029	.0017
SDev	.0046	.0006	.0002	.0010	.0226	.0000	.0019
%RSD	122.9	12.95	5.059	14.93	87.85	.9833	114.0
#1 #2	.0005	.0055	.0044	.0075	0098 0417	.0029	.0030
Elem Units Avge SDev %RSD	Sb206A ppm 0016 .0018 110.0	S_1820 ppm 28.63 .06 .2108	S_2203 ppm .0030 .0030 99.44	S_1960 ppm .0309 .0050 16.29	S_2068 ppm .0005 .0005 84.08	Ca3179 ppm 27.81 .02 .0635	Mg2790 ppm 20.49 .02
#1 #2	0028 0004	28.67 28.58	.0051	.0273	.0002	27.80 27.83	20.50
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0041	.0001	0289	.4780	0012	.0015	.0005
SDev	.0000	.0001	.0137	.0006	.0007	.0001	.0001
%RSD	.0711	142.1	47.29	.1326	56.14	5.657	11.78
#1 #2	.0041	.0002 0000	0386 0193	.4785 .4776	0007 0017	.0015	.0006
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0002	0331	1.641	45.02	1.122	.0001	.0332
SDev	.0008	.0008	.004	.14	.001	.0012	.0061
%RSD	351.7	2.295	.2534	.3022	.0866	1240.	18.34
#1	.0008	0326	1.644	45.11	1.123	.0009	.0289
#2	0003	0337	1.638	44.92	1.121	0007	.0375

MW-7-110419

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: _	Summit Env	vironmental Tech	Contract:	Jackpile	
Matrix: (so	il/water) _	water	Lab Sample ID: _	1107985-09	
Date Receiv	ed:	04/26/2011		·· ·	
% Solids: _			<u></u>		

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Q	М
7429-90-5	Aluminum	0.0286	J		Р
7440-36-0	Antimony	0.0030	J		P
7440-38-2	Arsenic	0.0028	J		Р
7440-39-3	Barium	0.0274	J		Р
7440-41-7	Beryllium	0.00072	U		P
7440-43-9	Cadmium	0.00082	U		Р
7440-70-2	Calcium	6.02		D	Р
7440-47-3	Chromium	0.00109	U		P
7440-48-4	Cobalt	0.0006	J		Р
7440-50-8	Copper	0.00183	Ū		P
7439-89-6	Iron	0.0174	J		P
7439-92-1	Lead	0.0029	J		P
7439-95-4	Magnesium	1.64			Р
7439-96-5	Manganese	0.00043	U		P
7440-02-0	Nickel	0.00116	U		Р
7440-09-7	Potassium	1.82			P
7782-49-2	Selenium	0.0296			P
7440-22-4	Silver	0.00039	U		Р
7440-23-5	Sodium	289		D	Р
7440-28-0	Thallium	0.0119	J		P
7440-62-2	Vanadium	0.00174	U	***************************************	P
7440-66-6	Zinc	0.0020	J		Р
7439-97-6	Mercury	0.00006	Ū		AV

Jomments	3:							
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Method: ZZJEM1_A Sample Name: 7985-09 Run Time: 04/29/11 18:36:38

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0070	.0003	.0002	.0006	.0274	.0028	0008
SDev	.0037	.0020	.0002	.0004	.0003	.0011	.0087
%RSD	52.37	603.6	99.23	64.26	1.179	39.64	1103.
#1 #2	.0096	.0018 0011	.0004	.0009	.0272 .0276	.0020	0069 .0053
Elem Units Avge SDev %RSD	Se196A ppm .0014 .0032 239.2	Cu3247 ppm 0013 .0004 27.20	Zn2138 ppm .0020 .0000	Ni2316 ppm .0006 .0002 34.24	Tl1908 ppm .0119 .0092 76.96	Be3130 ppm .0002 .0001 21.29	Sb2068 ppm .0076 .0012 15.94
#1 #2	.0037	0015 0010	.0020	.0007	.0054	.0002	.0084 .0067
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179 ppm 3.4020 .021 .6139	Mg2790
Units	ppm	ppm	ppm	ppm	ppm		ppm
Avge	0007	40.14	.0029	.0296	.0030		1.635
SDev	.0024	.18	.0025	.0031	.0012		.012
%RSD	330.0	.4595	87.64	10.61	38.45		.7184
#1 #2	0024 .0010	40.01	.0047	.0274 .0318	.0022	3.387 3.417	1.627 1.643
Elem	Mn2576	Co2286	Fe2714	A13082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0028	.0006	.0174	.0286	0143	.0046	.0007
SDev	.0001	.0003	.0080	.0031	.0002	.0006	.0000
%RSD	1.793	42.24	46.07	10.70	1.450	13.30	4.491
#1 #2	0028 0028	.0008	.0117	.0307	0144 0141	.0050	.0007 .0006
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0002	.2132	1.824	H423.4	4.298	.0110	.1221
SDev	.0003	.0027	.054	3.4	.029	.0021	.0002
%RSD	139.7	1.267	2.980	.7974	.6850	18.81	.1540
#1	.0004	.2113	1.786	H421.0	4.277	.0095	.1/219
#2		.2151	1.863	H425.8	4.319	.0125	/1222

Method: ZZJEM1_A Sample Name: 7985-09 x10 Operator: VVK Run Time: 05/11/11 12:14:12 Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0079 .0020 25.38	Pb220A ppm .0006 .0014 215.9	Cd2265 ppm .0001 .0003 230.5	Cr2677 ppm .0011 .0003 30.31	Ba4934 ppm .0034 .0000	As1890 ppm 0119 .0117 98.94	Se1960 ppm 0010 .0048 464.2
#1 #2	.0093 .0064	.0016 0003	0001 .0003	.0014	.0034	0036 0202	.0024 0045
Elem Units Avge SDev %RSD	Se196A ppm .0102 .0123 120.4	Cu3247 ppm .0050 .0000	Zn2138 ppm .0036 .0003 8.020	Ni2316 ppm .0046 .0017 36.00	Tl1908 ppm 0378 .0210 55.40	Be3130 ppm .0028 .0001 1.656	Sb2068 ppm .0042 .0010 23.27
#1 #2	.0015 .0189	.0050	.0038	.0034	0527 0230	.0028	.0049
Elem Units Avge SDev %RSD	Sb206A ppm 0018 .0037 207.4	S_1820 ppm 2.055 .010 .4903	S_2203 ppm .0033 .0016 47.11	S_1960 ppm .0337 .0066 19.62	S_2068 ppm .0012 .0028 229.2	Ca3179 ppm .6019 .0025 .4191	Mg2790 ppm .2548 .0000
#1 #2	.0008	2.048	.0044	.0290 .0384	.0032 0008	.6001 .6037	.2548 .2548
Elem Units Avge SDev %RSD	Mn2576 ppm .0010 .0000 .0488	Co2286 ppm 0004 .0006 141.4	Fe2714 ppm 0137 .0080 58.42	Al3082 ppm .4738 .0039 .8323	V_2924 ppm 0008 .0005 55.54	Mo2020 ppm .0022 .0012 55.00	Ti3349 ppm0000 .0002 565.6
#1 #2	.0010	0008 0000	0193 0080	.4766 .4711	0005 0011	.0013	.0001
Elem Units Avge SDev %RSD	Ag3280 ppm .0002 .0003 151.7	B_2496 ppm 0312 .0004 1.298	K_7664 ppm .5142 .0009 .1798	Na3302 ppm 28.89 .02 .0796	Si2881 ppm .6354 .0032 .5021	Sn1899 ppm .0009 .0000	P_1782 ppm .0344 .0058 16.87
#1 #2	0000 .0005	0315 0309	.5148 .5135	28.88 28.91	.6332 .6377	.0009	.0385

MW-8-110419

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit Env	ironmental Tech	Contract:	Jackpile	
Matrix: (soil/water)	water	_ Lab Sample ID: _	1107985-10	
Date Received:	04/26/2011	_		
% Solids:				
		_		

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Õ	М
7429-90-5	Aluminum	0.0227	J		Р
7440-36-0	Antimony	0.0096	J		Р
7440-38-2	Arsenic	0.0033	Ĵ		Р
7440-39-3	Barium	0.0389	J		Р
7440-41-7	Beryllium	0.00072	Ū		P
7440-43-9	Cadmium	0.00082	U		Р
7440-70-2	Calcium	15.3		D	Р
7440-47-3	Chromium	0.0012	J		P
7440-48-4	Cobalt	0.0010	J		P
7440-50-8	Copper	0.00183	Ū		Р
7439-89-6	Iron	0.0239	J		Р
7439-92-1	Lead	0.00210	U		Р
7439-95-4	Magnesium	1.59			Р
7439-96-5	Manganese	0.00043	U		P
7440-02-0	Nickel	0.0019	J		Р
7440-09-7	Potassium	5.59		N	Р
7782-49-2	Selenium	0.0317			P
7440-22-4	Silver	0.0011	J	N	Р
7440-23-5	Sodium	593		D	Р
7440-28-0	Thallium	0.00529	Ū		Р
7440-62-2	Vanadium	0.00174	Ū		P
7440-66-6	Zinc	0.00053	U		Р
7439-97-6	Mercury	0.00006	U		AV

J'omn	nents:							
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Method: ZZJEM1_A Sample Name: 7985-10 Run Time: 04/29/11 18:40:12

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm .0012 .0033 283.5	Pb220A ppm 0003 .0005 184.0	Cd2265 ppm .0001 .0000 11.89	Cr2677 ppm .0012 .0003 29.88	Ba4934 ppm .0389 .0002 .5535	As1890 ppm .0033 .0007 22.67	Se1960 ppm .0061 .0045 74.06
#1 #2	.0035 0012	.0001 0007	.0001	.0014	.0390	.0038	.0093
Elem Units Avge SDev %RSD	Se196A ppm .0037 .0035 95.34	Cu3247 ppm .0006 .0002 30.74	Zn2138 ppm 0016 .0001 3.069	Ni2316 ppm .0019 .0008 41.70	Tl1908 ppm .0027 .0041 153.2	Be3130 ppm .0001 .0000 39.05	Sb2068 ppm .0053 .0012 22.66
#1 #2	.0062 .0012	.0004	0016 0017	.0025 .0014	0002 .0056	.0001	.0062
Elem Units Avge SDev %RSD	Sb206A ppm .0102 .0001 .5087	S_1820 ppm H147.0 .1 .0351	S_2203 ppm .0004 .0015 332.0	S_1960 ppm .0317 .0039 12.13	S_2068 ppm .0096 .0004 3.856	Ca3179 ppm 13.20 .01 .0947	Mg2790 ppm 1.594 .007 .4550
#1 #2	.0102	H147.0 H147.0	.0015 0006	.0345	.0098	13.21 13.19	1.599 1.589
Elem Units Avge SDev %RSD	Mn2576 ppm 0173 .0000	Co2286 ppm .0010 .0001 8.362	Fe2714 ppm .0239 .0049 20.58	A13082 ppm .0227 .0030 13.32	V_2924 ppm 0298 .0006 2.074	Mo2020 ppm .0223 .0011 4.736	Ti3349 ppm .0001 .0003 271.1
#1 #2	0172 0173	.0010	.0274	.0248 .0206	0293 0302	.0216 .0231	.0004
Elem Units Avge SDev %RSD	Ag3280 ppm .0011 .0000 1.788	B_2496 ppm .7883 .0001 .0172	K_7664 ppm 5.590 .001 .0139	Na3302 ppm H882.7 .7 .0849	Si2881 ppm 5.009 .001 .0185	Sn1899 ppm .0104 .0021 20.41	P_1782 ppm .0818 .0016 1.998
#1 #2	.0011	.7884 .7882	5.591 5.590	H883.3 H882.2	5.009 5.010	.0119	.0807

Method: ZZJEM1_A Sample Name: 7985-10 x10 Operator: VVK Run Time: 05/11/11 12:17:46 Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem	Pb2203	Pb220A	Cd2265	Cr2677 ppm .0007 .0009 125.7	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm		ppm	ppm	ppm
Avge	0032	0009	0002		.0046	0111	.0030
SDev	.0239	.0035	.0002		.0001	.0028	.0026
%RSD	740.5	383.9	141.8		1.589	25.17	89.04
#1 #2	.0137	.0016 0034	.0000	.0001	.0045	0091 0130	.0011
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0011	.0048	.0018	.0050	.0059	.0027	.0024
SDev	.0000	.0006	.0005	.0010	.0221	.0000	.0037
%RSD	2.525	12.39	24.86	20.20	377.1	.7570	154.0
#1 #2	0011 0010	.0053	.0022 .0015	.0043	0098 .0215	.0028	.0050
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0015	7.230	0013	.0258	.0028	1.533	.2133
SDev	.0002	.019	.0103	.0015	.0012	.013	.0046
%RSD	10.88	.2656	789.8	5.890	43.66	.8227	2.168
#1 #2	.0014	7.216 7.243	.0060 0086	.0269 .0247	.0036 .0019	1.524 1.542	.2166
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0001	0002	0094	.4813	0023	.0017	.0000
SDev	.0001	.0002	.0084	.0198	.0002	.0002	.0001
%RSD	50.06	141.4	88.97	4.113	9.312	10.10	209400.
#1	0002	0003	0035	.4953	0022	.0018	.0000
#2	0001	.0000	0153	.4673	0025	.0016	
Elem Units Avge SDev %RSD	Ag3280 ppm .0002 .0001 22.91	B_2496 ppm .0405 .0000	K_7664 ppm .6993 .0000	Na3302 ppm 59.30 .29 .4943	Si2881 ppm .7232 .0008 .1151	Sn1899 ppm .0003 .0001 42.94	P_1782 ppm .0262 .0054 20.64
#1 #2	.0002	.0405	.6992 .6993	59.51 59.09	.7226 .7238	.0002	.0301

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name:	Summit Env	ironmental Tech	Contract:	Jackpile
Matrix: (s	oil/water)	water	Lab Sample ID:	1107985-11
Date Recei	ved:	04/26/2011		
% Solids:	- Andrea			
Concentrat	ion Units (mg	g/L or mg/kg dry we	eight): mg/L	

CAS No.	Analyte	Concentration	C	Q	М
7429-90-5	Aluminum	0.0105	U		Р
7440-36-0	Antimony	0.0074	J		P
7440-38-2	Arsenic	0.0102	J		P
7440-39-3	Barium	0.0132	J		Р
7440-41-7	Beryllium	0.00072	Ū		P
7440-43-9	Cadmium	0.00082	U		P
7440-70-2	Calcium	214		D	P
7440-47-3	Chromium	0.0014	J		P
7440-48-4	Cobalt	0.0017	J		P
7440-50-8	Copper	0.00183	U		P
7439-89-6	Iron	1.61			P
7439-92-1	Lead	0.00210	Ū		P
7439-95-4	Magnesium	252		D	P
7439-96-5	Manganese	0.693			P
7440-02-0	Nickel	0.0057	J		Р
7440-09-7	Potassium	13.3			P
7782-49-2	Selenium	0.0395			P
7440-22-4	Silver	0.00039	Ū		P
7440-23-5	Sodium	1410		D	P
7440-28-0	Thallium	0.0099	J		P
7440-62-2	Vanadium	0.00174	U		P
7440-66-6	Zinc	0.00053	Ū		P
7439-97-6	Mercury	0.00006			AV

Com	ments:						
		 	70.00	 	7000		
		 		 ~ <u>~</u>	97	-10.	

Method: ZZJEM1_A Sample Name: 7985-11 Run Time: 04/29/11 18:50:54

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm .0013 .0024 184.8	Pb220A ppm .0009 .0021 240.0	Cd2265 ppm .0005 .0000 9.707	Cr2677 ppm .0014 .0004 27.60	Ba4934 ppm .0132 .0003 2.452	As1890 ppm .0102 .0025 24.01	Se1960 ppm .0089 .0059 66.12
#1 #2	.0030	0006 .0024	.0004	.0012 .0017	.0129 .0134	.0085 .0120	.0131
Elem Units Avge SDev %RSD	Se196A ppm .0080 .0008 10.05	Cu3247 ppm 0020 .0014 69.82	Zn2138 ppm 0243 .0004 1.837	Ni2316 ppm .0057 .0012 20.99	Tl1908 ppm .0099 .0161 162.0	Be3130 ppm .0004 .0000 1.234	Sb2068 ppm .0035 .0057 162.3
#1 #2	.0086	0029 0010	0246 0240	.0065 .0048	0014 .0213	.0004	0005 .0076
Elem Units Avge SDev %RSD	Sb206A ppm .0078 .0063 80.57	S_1820 ppm H1213. 3. .2622	S_2203 ppm .0012 .0006 52.83	S_1960 ppm .0395 .0001 .2074	S_2068 ppm .0074 .0022 29.38	Ca3179 ppm H191.7 .7	Mg2790 ppm H216.3 1.0 .4595
#1 #2	.0122	H1211. H1216.	.0007 .0016	.0395	.0089 .0058	H191.3 H192.2	H215.6 H217.0
Elem Units Avge SDev %RSD	Mn2576 ppm .6927 .0020 .2916	Co2286 ppm .0017 .0001 7.515	Fe2714 ppm 1.605 .007 .4117	Al3082 ppm .0070 .0081 115.6	V_2924 ppm 1255 .0004 .2946	Mo2020 ppm .0142 .0006 4.206	Ti3349 ppm 0010 .0003 26.71
#1 #2	.6913 .6942	.0016	1.600 1.609	.0013 .0128	1258 1252	.0137	0011 0008
Elem Units Avge SDev %RSD	Ag3280 ppm .0003 .0005 136.8	B_2496 ppm .5425 .0030 .5518	K_7664 ppm 13.32 .03 .2335	Na3302 ppm H3679. 5.	Si2881 ppm 9.354 .023 .2458	Sn1899 ppm .0020 .0039 193.4	P_1782 ppm .0574 .0017 2.945
#1 #2	.0000	.5404 .5446	13.34 13.30	H3676. H3683.	9.337 9.370	0007 .0048	.0562 .0586

Method: ZZJEM1_A Sample Name: 7985-11 x10 Operator: VVK

Run Time: 05/11/11 12:41:55

Elem Units Avge SDev %RSD	Pb2203 ppm .0063 .0079 124.1	Pb220A ppm .0008 .0016 206.7	Cd2265 ppm .0010 .0002 22.12	Cr2677 ppm .0013 .0001 7.904	Ba4934 ppm .0026 .0001 2.668	As1890 ppm 0009 .0015 154.5	Se1960 ppm .0110 .0019 17.70
#1 #2	.0008	0004 .0019	.0012	.0014	.0026 .0025	.0001 0020	.0096 .0124
Elem Units Avge SDev %RSD	Se196A ppm .0084 .0013 14.82	Cu3247 ppm .0070 .0004 6.061	Zn2138 ppm .0046 .0000	Ni2316 ppm .0057 .0019 32.78	Tl1908 ppm 0148 .0428 289.1	Be3130 ppm .0025 .0001 2.663	Sb2068 ppm .0130 .0020 15.59
#1 #2	.0093 .0075	.0073	.0046	.0044	0451 .0155	.0025	.0144
Elem Units Avge SDev %RSD	Sb206A ppm .0051 .0004 7.603	S_1820 ppm 71.01 .02 .0258	S_2203 ppm .0029 .0037 126.9	S_1960 ppm .0367 .0002 .4254	S_2068 ppm .0087 .0009 10.70	Ca3179 ppm 21.37 .06 .2674	Mg2790 ppm 25.21 .08 .3019
#1 #2	.0054 .0048	71.02 71.00	.0003 .0055	.0368 .0366	.0094	21.33	25.26 25.16
Elem Units Avge SDev %RSD	Mn2576 ppm .0900 .0000	Co2286 ppm .0011 .0005 47.60	Fe2714 ppm .1411 .0202 14.29	Al3082 ppm .3174 .0008 .2601	V_2924 ppm 0090 .0007 7.556	Mo2020 ppm .0065 .0009 14.14	Ti3349 ppm .0015 .0004 26.05
#1 #2	.0901 .0900	.0007	.1268 .1553	.3168 .3180	0095 0085	.0072 .0059	.0018 .0012
Elem Units Avge SDev %RSD	Ag3280 ppm .0022 .0005 25.04	B_2496 ppm .0310 .0005 1.670	K_7664 ppm 1.771 .004 .2054	Na3302 ppm H291.9 1.7 .5847	Si2881 ppm 1.189 .006 .4718	Sn1899 ppm .0005 .0001 9.094	P_1782 ppm .0403 .0034 8.492
#1 #2	.0026	.0313	1.774 1.768	H293.2 H290.7	1.193 1.185	.0005	.0379

Method: ZZJEM1_A Sample Name: 7985-11 x50 Operator: VVK Run Time: 05/11/11 15:48:58

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0052	.0015	.0002	.0007	.0007	0063	0067
SDev	.0111	.0011	.0001	.0001	.0001	.0085	.0051
%RSD	213.9	74.08	35.33	21.81	9.428	135.1	76.25
#1 #2	.0130	.0022	.0002	.0008	.0007	0003 0123	0031 0103
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908 ppm0142 .0104 73.43	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm		ppm	ppm
Avge	.0037	.0037	.0423	.0115		.0011	.0019
SDev	.0032	.0003	.0040	.0009		.0000	.0010
%RSD	87.80	7.379	9.556	7.739		1.040	55.34
#1 #2	.0059 .0014	.0035	.0394 .0451	.0121	0216 0068	.0011	.0012 .0026
Elem Units Avge SDev %RSD	Sb206A ppm 0050 .0029 58.70	S_1820 ppm 17.11 .15 .8493	S_2203 ppm .0030 .0044 149.3	S_1960 ppm .0275 .0039 14.04	S_2068 ppm 0017 .0023 136.8	Ca3179 ppm 4.865 .002	Mg2790 ppm 4.951 .030 .5962
#1	0070	17.21	.0061	.0302	0033	4.864	4.972
#2	0029	17.01	0002		0001	4.866	4.930
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0196	.0001	.0176	.2644	0012	.0009	.0005
SDev	.0001	.0001	.0066	.0006	.0002	.0014	.0001
%RSD	.2687	48.22	37.26	.2087	14.97	154.3	16.64
#1 #2	.0196 .0197	.0001	.0223 .0130	.2640 .2648	0013 0011	0001 .0019	.0006
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0000	0111	1.120	28.25	.2568	.0005	.0799
SDev	.0003	.0037	.079	.42	.0007	.0005	.0042
%RSD	275800.	33.21	7.025	1.484	.2878	101.0	5.288
#1 #2	.0002	0085 0137	1.065 1.176	28.54 27.95	.2573 .2562	.0008	.0770 .0829

MW-MD-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit En	vironmental Tech	Contract:	Jackpile	
Matrix: (soil/water)	water	_ Lab Sample ID: _	1107985-12	
Date Received:	04/26/2011	_		
% Solids:	- W	_		

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Q	M
7429-90-5	Aluminum	0.0105	U		Р
7440-36-0	Antimony	0.0041	J		Р
7440-38-2	Arsenic	0.0064	J		P
7440-39-3	Barium	0.0116	J		Р
7440-41-7	Beryllium	0.00072	U		P
7440-43-9	Cadmium	0.00082	U		P
7440-70-2	Calcium	200		D	P
7440-47-3	Chromium	0.00109	U		P
7440-48-4	Cobalt	0.0013	J		Р
7440-50-8	Copper	0.00183	U		P
7439-89-6	Iron	1.45			Р
7439-92-1	Lead	0.00210	Ū		P
7439-95-4	Magnesium	237		D	P
7439-96-5	Manganese	0.654			P
7440-02-0	Nickel	0.0048	J		Р
7440-09-7	Potassium	12.6			Р
7782-49-2	Selenium	0.0271			Р
7440-22-4	Silver	0.00039	U		P
7440-23-5	Sodium	1310		D	Р
7440-28-0	Thallium	0.00529	Ū		P
7440-62-2	Vanadium	0.00174	U		Р
7440-66-6	Zinc	0.00053	Ū		P
7439-97-6	Mercury	0.00006	U		AV

Jomments:					
	- My		 	7%.	
		Metro	 	100	

Method: ZZJEM1_A Sample Name: 7985-12 Run Time: 04/29/11 18:54:28

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm .0096 .0052 54.23	Pb220A ppm 0022 .0018 80.18	Cd2265 ppm .0004 .0001 22.28	Cr2677 ppm .0008 .0001 13.06	Ba4934 ppm .0116 .0001 .9243	As1890 ppm .0064 .0030 46.44	Se1960 ppm .0110 .0006 4.965
#1 #2	.0133	0010 0035	.0004	.0007	.0117	.0086	.0106 .0114
Elem Units Avge SDev %RSD	Se196A ppm .0014 .0023 162.5	Cu3247 ppm0020 .0006 29.68	Zn2138 ppm 0262 .0003 1.166	Ni2316 ppm .0048 .0002 3.297	Tl1908 ppm .0049 .0032 65.15	Be3130 ppm .0002 .0000 2.015	Sb2068 ppm 0024 .0089 364.6
#1 #2	.0030	0024 0016	0264 0260	.0047	.0072 .0027	.0002	.0038 0087
Elem Units Avge SDev %RSD	Sb206A ppm .0062 .0068 109.7	S_1820 ppm H1169. 6.	S_2203 ppm .0018 .0029 167.2	S_1960 ppm .0271 .0014 5.021	S_2068 ppm .0041 .0016 38.21	Ca3179 ppm H182.5 1.1 .5797	Mg2790 ppm H205.4 1.1 .5237
#1 #2	.0014	H1174. H1165.	.0038	.0280 .0261	.0030	H183.3 H181.8	H206.2 H204.6
Elem Units Avge SDev %RSD	Mn2576 ppm .6540 .0044 .6676	Co2286 ppm .0013 .0001 7.001	Fe2714 ppm 1.449 .008 .5507	Al3082 ppm 0015 .0066 445.6	V_2924 ppm 1192 .0004 .3510	Mo2020 ppm .0121 .0001 .9847	Ti3349 ppm 0014 .0001 10.96
#1 #2	.6571 .6510	.0012	1.455 1.444	.0032 0062	1195 1189	.0122 .0120	0013 0015
Elem Units Avge SDev %RSD	Ag3280 ppm 0000 .0009 2067.	B_2496 ppm .5093 .0046 .8991	K_7664 ppm 12.61 .02 .1389	Na3302 ppm \ H3508. 16. .4420	Si2881 ppm 8.920 .060 .6709	Sn1899 ppm .0007 .0013 190.6	P_1782 ppm .0539 .0042 7.733
#1 #2	0007 .0006	.5126 .5061	12.63 12.60	H3519. H3497.	8.962 8.878	.0016 0002	.0569 .0510

Method: ZZJEM1_A Sample Name: 7985-12 x10 Operator: VVK Run Time: 05/11/11 12:45:30

Elem Units Avge SDev %RSD	Pb2203 ppm 0064 .0011 17.62	Pb220A ppm .0026 .0003 9.867	Cd2265 ppm .0006 .0004 62.32	Cr2677 ppm .0013 .0001 7.904	Ba4934 ppm .0021 .0001 3.289	As1890 ppm 0064 .0058 90.51	Se1960 ppm .0101 .0117 115.7
#1 #2	0056 0072	.0028	.0009	.0012	.0021	0023 0105	.0184
Elem Units Avge SDev %RSD	Se196A ppm .0066 .0033 50.11	Cu3247 ppm .0079 .0001 .5992	Zn2138 ppm .0133 .0011 8.534	Ni2316 ppm .0104 .0015 14.45	Tl1908 ppm 0079 .0154 194.5	Be3130 ppm .0021 .0001 4.390	Sb2068 ppm .0049 .0034 70.00
#1 #2	.0090	.0078 .0079	.0125 .0141	.0093 .0114	0188 .0030	.0022	.0073
Elem Units Avge SDev %RSD	Sb206A ppm .0013 .0013 104.5	S_1820 ppm 67.42 .20 .3014	S_2203 ppm 0001 .0005 925.6	S_1960 ppm .0369 .0037 10.05	S_2068 ppm .0034 .0002 5.096	Ca3179 ppm 19.95 .07	Mg2790 ppm 23.69 .11 .4630
#1 #2	.0003	67.57 67.28	.0003	.0395 .0343	.0035	20.00	23.77 23.61
Elem Units Avge SDev %RSD	Mn2576 ppm .0834 .0006 .6966	Co2286 ppm .0012 .0006 47.53	Fe2714 ppm .1270 .0016 1.303	Al3082 ppm .3469 .0031 .9040	V_2924 ppm 0082 .0003 4.110	Mo2020 ppm .0024 .0023 96.42	Ti3349 ppm .0008 .0002 21.21
#1 #2	.0838 .0830	.0008	.1258 .1282	.3491 .3446	0079 0084	.0040	.0009 .0007
Elem Units Avge SDev %RSD	Ag3280 ppm .0010 .0001 5.310	B_2496 ppm .0224 .0016 7.312	K_7664 ppm 2.154 .001 .0239	Na3302 ppm H271.5 2.3 .8643	Si2881 ppm 1.118 .008 .7413	Sn1899 ppm 0003 .0000 7.318	P_1782 ppm .0321 .0051 15.99
#1 #2	.0010	.0236 .0213	2.153 2.154	H273.2 H269.9	1.123 1.112	0003 0003	.0357

Method: ZZJEM1_A Sample Name: 7985-12 x50 Operator: VVK Run Time: 05/11/11 15:53:59 Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

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Elem Units Avge SDev %RSD	Pb2203 ppm 0072 .0011 14.95	Pb220A ppm .0016 .0020 121.2	Cd2265 ppm .0002 .0003 186.0	Cr2677 ppm .0013 .0001 5.898	Ba4934 ppm .0006 .0001 20.20	As1890 ppm 0069 .0035 50.97	Se1960 ppm 0038 .0028 72.94
#1 #2	0064 0079	.0002	.0004	.0012 .0013	.0005	0044 0094	0058 0018
Elem Units Avge SDev %RSD	Se196A ppm 0003 .0004 104.8	Cu3247 ppm .0033 .0002 5.387	Zn2138 ppm .0273 .0051 18.77	Ni2316 ppm .0082 .0008 10.35	Tl1908 ppm 0079 .0000	Be3130 ppm .0011 .0000 2.603	Sb2068 ppm .0001 .0006 518.5
#1 #2	0006 0001	.0035	.0236 .0309	.0088 .0076	0079 0079	.0012	0003 .0006
Elem Units Avge SDev %RSD	Sb206A ppm 0028 .0005 18.86	S_1820 ppm 16.06 .01 .0479	S_2203 ppm 0010 .0009 97.92	S_1960 ppm .0258 .0012 4.607	S_2068 ppm 0008 .0001 6.518	Ca3179 ppm 4.296 .067 1.567	Mg2790 ppm 4.630 .001
#1 #2	0024 0032	16.07 16.06	0016 0003	.0249 .0266	0007 0008	4.249	4.629 4.630
Elem Units Avge SDev %RSD	Mn2576 ppm .0183 .0003 1.660	Co2286 ppm .0004 .0002 47.38	Fe2714 ppm .0181 .0137 75.83	Al3082 ppm .2676 .0050 1.873	V_2924 ppm 0008 .0003 41.81	Mo2020 ppm 0007 .0012 174.7	Ti3349 ppm .0008 .0006 70.71
#1 #2	.0180 .0185	.0003	.0084	.2640 .2711	0010 0005	.0002 0016	.0004
Elem Units Avge SDev %RSD	Ag3280 ppm .0002 .0001 55.41	B_2496 ppm 0229 .0001 .5790	K_7664 ppm .7258 .0725 9.993	Na3302 ppm 26.10 .18 .6738	Si2881 ppm .2412 .0020 .8314	Sn1899 ppm .0007 .0001 18.62	P_1782 ppm .0413 .0117 28.31
#1 #2	.0001	0228 0230	.6745 .7771	25.98 26.23	.2426 .2398	.0006 .0008	.0330

NPOP20E-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Sun	mmit Environmental Tech	Contract:	Jackpile	
Matrix: (soil/w	water) water	_ Lab Sample ID: _	1107985-13	
Date Received:	04/26/2011	_		
% Solids:		_		
Concentration I	Jnits (mg/L or mg/kg dry we:	ich+\ ma/I		
Concentration	mires (mg/r or mg/kg dry we-	rgiri): _ mg/ h		

CAS No.	Analyte	Concentration	С	Q	М
7429-90-5	Aluminum	0.822			Р
7440-36-0	Antimony	0.00237	Ū		P
7440-38-2	Arsenic	0.00206	U		Р
7440-39-3	Barium	0.0112	J		P
7440-41-7	Beryllium	0.0067			P
7440-43-9	Cadmium	0.00082	U		P
7440-70-2	Calcium	571		D	Р
7440-47-3	Chromium	0.0604			P
7440-48-4	Cobalt	0.0692	J		Р
7440-50-8	Copper	0.00183	Ū		P
7439-89-6	Iron	3.18			Р
7439-92-1	Lead	0.0519			Р
7439-95-4	Magnesium	276		D	Р
7439-96-5	Manganese	4.633			Р
7440-02-0	Nickel	0.0374			Р
7440-09-7	Potassium	39.7			Р
7782-49-2	Selenium	0.0309			Р
7440-22-4	Silver	0.0522			Р
7440-23-5	Sodium	1260		D	Р
7440-28-0	Thallium	0.00529	U		Р
7440-62-2	Vanadium	0.00174	U		Р
7440-66-6	Zinc	0.0107	J		Р
7439-97-6	Mercury	0.00006	Ŭ		AV

Com	ments:							
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Method: ZZJEM1_A Sample Name: 7985-13 Run Time: 04/29/11 18:58:01

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm 0120 .0065 53.84	Pb220A ppm .0841 .0019 2.204	Cd2265 ppm .0003 .0002 51.91	Cr2677 ppm .0604 .0006 1.024	Ba4934 ppm .0112 .0002 1.448	As1890 ppm 0007 .0001 20.05	Se1960 ppm .0321 .0002 .6024
#1 #2	0166 0074	.0854	.0005	.0608 .0600	.0113	0008 0006	.0322
Elem Units Avge SDev %RSD	Se196A ppm 0198 .0020	Cu3247 ppm 0254 .0011 4.168	Zn2138 ppm .0107 .0004 3.983	Ni2316 ppm .0374 .0001 .3332	Tl1908 ppm .0037 .0020 55.71	Be3130 ppm .0067 .0000 .4195	Sb2068 ppm 0472 .0013 2.705
#1 #2	0184 0212	0246 0261	.0110	.0373 .0375	.0051	.0067 .0067	0463 0481
Elem Units Avge SDev %RSD	Sb206A ppm 1657 .0033 1.971	S_1820 ppm H1033. 6. .5438	S_2203 ppm .0519 .0010 1.858	S_1960 ppm .0309 .0010 3.232	S_2068 ppm 1252 .0026 2.079	Ca3179 ppm H509.4 2.4 .4784	Mg2790 ppm H234.7 1.3 .5450
#1 #2	1634 1680	H1029. H1037.	.0512	.0302	1234 1271	H507.6 H511.1	H233.8 H235.6
Elem Units Avge SDev %RSD	Mn2576 ppm 4.633 .024 .5189	Co2286 ppm .0692 .0004 .6516	Fe2714 ppm 3.181 .007 .2351	Al3082 ppm .8217 .0013 .1577	V_2924 ppm 0829 .0007 .8222	Mo2020 ppm .0076 .0017 22.85	Ti3349 ppm .0085 .0005 5.958
#1 #2	4.616 4.650	.0695 .0689	3.186 3.175	.8208 .8226	0824 0834	.0089	.0089
Elem Units Avge SDev %RSD	Ag3280 ppm .0522 .0012 2.373	B_2496 ppm .1585 .0015 .9579	K_7664 ppm 39.68 .23	Na3302 ppm H1692. 11. .6277	Si2881 ppm 5.300 .025 .4728	Sn1899 ppm 0065 .0002 2.936	P_1782 ppm .0744 .0034 4.598
#1 #2	.0531 .0514	.1596 .1574	39.52 39.85	H1685. H1700.	5.282 5.318	0064 0066	.0768 .0720

Method: ZZJEM1_A Sample Name: 7985-13 x10 Operator: VVK

Run Time: $05/1\overline{1}/11$ 12:49:04

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0218	.0233	.0006	.0208	.0018	0057	.0070
SDev	.0136	.0017	.0002	.0005	.0001	.0033	.0045
%RSD	62.46	7.154	32.23	2.464	3.822	58.50	64.31
#1 #2	0315 0122	.0221	.0004	.0204 .0212	.0019 .0018	0033 0081	.0038
Elem Units Avge SDev %RSD	Se196A ppm 0035 .0064 182.2	Cu3247 ppm .0000 .0001 282.8	Zn2138 ppm .0035 .0005	Ni2316 ppm .0101 .0022 22.16	Tl1908 ppm 0023 .0279 1212.	Be3130 ppm .0037 .0000 .9175	Sb2068 ppm .0056 .0000
#1 #2	0080 .0010	0000 .0001	.0032	.0117 .0085	0221 .0174	.0037	.0056
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0137	61.09	.0085	.0308	0063	57.08	27.63
SDev	.0022	.74	.0056	.0058	.0015	.52	.41
%RSD	15.84	1.214	66.29	18.71	24.27	.9152	1.498
#1 #2	0122 0152	61.61 60.56	.0045	.0267	0052 0074	57.45 56.71	27.93 27.34
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.5870	.0101	.1020	.6028	0085	.0009	.0026
SDev	.0058	.0004	.0004	.0040	.0003	.0008	.0001
%RSD	.9882	3.927	.4093	.6709	3.911	91.51	2.143
#1	.5911	.0098	.1017	.6057	0083	.0003	.0025
#2	.5829	.0103		.5999	0088	.0015	.0026
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0073	0189	3.051	H126.4	.6928	0005	.0208
SDev	.0005	.0016	.021	2.4	.0087	.0002	.0026
%RSD	7.078	8.585	.6918	1.896	1.251	46.37	12.59
#1	.0076	0178	3.065	H128.1	.6989	0007	.0189
#2	.0069	0201	3.036	H124.8	.6866	0003	.0226

PR-SED-01-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name:	Summit Env	ironmental Tech	Contract:	Jackpile	
Matrix: (so	oil/water) _	soil	Lab Sample ID: _	1107985-14	
Date Recei	ved:	04/26/2011			
% Solids:	**************************************				

Concentration Units (mg/L or mg/kg dry weight): mg/kg

CAS No.	Analyte	Concentration	С	Q	М
7429-90-5	Aluminum	31900		D	P
7440-36-0	Antimony	1.88	J		Р
7440-38-2	Arsenic	7.82			Р
7440-39-3	Barium	187			Р
7440-41-7	Beryllium	1.19	J		P
7440-43-9	Cadmium	0.901			Р
7440-70-2	Calcium	74200		D	P
7440-47-3	Chromium	17.7			Р
7440-48-4	Cobalt	7.30			P
7440-50-8	Copper	18.9			P
7439-89-6	Iron	27300		D	P
7439-92-1	Lead	9.69	J		Р
7439-95-4	Magnesium	12300		D	Р
7439-96-5	Manganese	558			Р
7440-02-0	Nickel	14.7			P
7440-09-7	Potassium	9600		D	Р
7782-49-2	Selenium	0.158	U		Р
7440-22-4	Silver	0.158	Ū		Р
7440-23-5	Sodium	5610			P
7440-28-0	Thallium	0.253	U		Р
7440-62-2	Vanadium	44.8			Р
7440-66-6	Zinc	65.0			Р
7439-97-6	Mercury	0.0632	U		AV

Comm	ents:								
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Method: ZZJEM1 A Sample Name: 7985-14s

Run Time: 05/12/11 12:37:58

Comment: 6010/200.7 CPI 4400-013 Mode: CONC Corr. Factor: 1 lot 10B101-

Blem	•					v			
#2		Units Avge SDev	ppm .2246 .0116	ppm .1404 .0026	ppm .0114 0 .0003	ppm .2243 .0033	ppm 2.364 .017	ppm .0985 .0117	ppm .5500 .0211
Units ppm									
Elem Sb206A S 1820 S 2203 S 1960 S 2068 Ca3179 Mg2790 ppm ppm ppm ppm ppm ppm ppm ppm ppm pp		Units Avge SDev	ppm <-1.000 .031	ppm .2395 0	ppm 5 .8223 6 .0103	ppm .1866 /// .0049	ppm 0565(1) .0026	ppm .0151	ppm .0267 .0013
Units ppm									
#2 .0135 67.98 .12627511 .0190 758.5 135.4 Elem Mn2576 C02286 ppm ppm 290.0 387.2 .5670 .0038 1.024 SDev .090 .0012 3.6 3.1 .0070 .0009 .010 %RSD 1.268 1.304 1.252 .7890 1.229 24.84 .9877 #1 7.003 .0916 287.5 385.0 .5620 .0031 1.017 #2 7.130 .0933 292.6 389.4 .5719 .0044 1.031 Elem Ag3280 B 2496 K 7664 Na3302 Si2881 Sn1899 ppm ppm ppm ppm ppm ppm ppm ppm ppm p		Units Avge SDev	ppm .0204 .0098	ppm 67.45 .75	ppm .1226 .0050	ppm 7482 .0041	.0238 .0068	ppm 752.10 9.1	ppm 134.3 1.5
Units ppm									
#2 7.130 .0933 292.6 389.4 .5719 .0044 1.031 Elem Ag3280 B_2496 K_7664 Na3302 Si2881 Sn1899 P_1782 Units ppm ppm ppm ppm ppm ppm ppm ppm ppm pp		Units Avge SDev	ppm 7.066 .090	ppm .0924/b	ppm 290.00	ppm 387.2 3.1	ppm W.5670 .0070	ppm .0038 .0009	ppm 1.024 .010
Units ppm ppm ppm ppm ppm ppm ppm ppm ppm pp									
u =		Units Avge SDev	ppm 0008 0007	ppm .7143 .0065	ppm 141.5	ppm 70.95 (b)	ppm 9.046 .092	ppm .0168 .0001	ppm 7.163 .092

Method: ZZJEM1_A Sample Name: 7985-14s x10 Operator: VVK Run Time: $05/1\overline{2}/11$ 12:42:39

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0273	.0230	.0016	.0266	.2445	.0153	.0633
SDev	.0196	.0019	.0002	.0007	.0010	.0006	.0061
%RSD	72.07	8.360	12.02	2.467	.4035	4.265	9.692
#1 #2	.0411	.0244	.0015	.0270 .0261	.2452	.0148 .0157	.0676 .0590
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	2210	.0316	.1201	.0389	0028	.0105	.0023
SDev	.0038	.0012	.0012	.0003	.0171	.0017	.0011
%RSD	1.698	3.915	.9900	.6529	612.8	15.78	48.14
#1 #2	2236 2183	.0307	.1193 .1210	.0387	.0093 0149	.0093 .0117	.0031 .0015
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790 ppm 15.54 0 .08 .5213
Units	ppm	ppm	ppm	ppm	ppm	ppm	
Avge	0022	8.810	.0192	0529	.0003	93.83	
SDev	.0012	.020	.0078	.0022	.0012	.91	
%RSD	53.03	.2321	40.56	4.241	483.8	.9682	
#1 #2	0014 0031	8.824 8.795	.0247	0513 0545	.0011	94.47 93.18	15.60 15.48
Elem	Mn2576	Co2286 ppm .0093 .0000 .0090	Fe2714	A13082 (F	V_2924	Mo2020	Ti3349
Units	ppm		ppm	ppm	ppm	ppm	ppm
Avge	.8466		34.50	40.36 7	.0636	0002	.1126
SDev	.0070		.21	.24	.0002	.0015	.0008
%RSD	.8229		.6129	.5851	.3222	742.5	.6767
#1	.8516	.0093	34.65	40.52	.0635	.0009	.1132
#2	.8417	.0093	34.35	40.19	.0638	0013	.1121
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0007	.1038	12.14	5.705	1.082	.0030	1.013
SDev	.0001	.0016	.04	.157	.006	.0002	.007
%RSD	15.50	1.507	.3517	2.757	.5209	6.826	.7405
#1 #2	0006 0008	.1049	12.11 12.17	5.817 5.594	1.086 1.078	.0032	1.008

PR-SW-01-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit Env	ironmental Tech	Contract:	Jackpile
Matrix: (soil/water) _	water	Lab Sample ID:	1107985-15
Date Received:	04/26/2011		
% Solids:		_	
Concentration Units (mg	g/L or mg/kg dry we	eight): _mg/L	

CAS No.	Analyte	Concentration	C	Q	M
7429-90-5	Aluminum	0.0105	U		Р
7440-36-0	Antimony	0.0033	J		P
7440-38-2	Arsenic	0.0075	J		Р
7440-39-3	Barium	0.140			P
7440-41-7	Beryllium	0.00072	U		P
7440-43-9	Cadmium	0.00082	U		P
7440-70-2	Calcium	1010		D	P
7440-47-3	Chromium	0.00109	Ū		P
7440-48-4	Cobalt	0.0009	J		Р
7440-50-8	Copper	0.00183	U		Р
7439-89-6	Iron	0.269			Р
7439-92-1	Lead	0.00210	U		Р
7439-95-4	Magnesium	471		D	P
7439-96-5	Manganese	0.970			P
7440-02-0	Nickel	0.0032	J		P
7440-09-7	Potassium	92.8		D	Р
7782-49-2	Selenium	0.0302			P
7440-22-4	Silver	0.00039	U		P
7440-23-5	Sodium	1200		D	P
7440-28-0	Thallium	0.0130	J		Р
7440-62-2	Vanadium	0.00174	Ū		Р
7440-66-6	Zinc	0.00053	Ū		P
7439-97-6	Mercury	0.00006	U		AV
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Comm	ents:							
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Method: ZZJEM1_A Sample Name: 7985-15 Run Time: 04/29/11 19:01:35

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm 0043 .0084 195.7	Pb220A ppm 0050 .0013 26.15	Cd2265 ppm .0002 .0000 13.66	Cr2677 ppm .0000 .0001 1233.	Ba4934 ppm .0140 .0001 .7686	As1890 ppm .0075 .0014 18.15	Se1960 ppm .0125 .0090 72.08
#1 #2	0102 .0016	0059 0041	.0003	.0001	.0139	.0085	.0189 .0061
Elem Units Avge SDev %RSD	Se196A ppm 0024 .0021 84.79	Cu3247 ppm 0036 .0002 5.812	Zn2138 ppm 0063 .0000 .4594	Ni2316 ppm .0032 .0007 23.09	Tl1908 ppm .0130 .0082 62.85	Be3130 ppm 0000 .0000 10990.	Sb2068 ppm0088 .0037 42.33
#1 #2	0039 0010	0038 0035	0063 0064	.0027	.0072 .0188	.0000	0115 0062
Elem Units Avge SDev %RSD	Sb206A ppm .0079 .0041 51.52	S_1820 ppm H1484. 1.	S_2203 ppm 0043 .0037 85.63	S_1960 ppm .0302 .0016 5.419	S_2068 ppm .0033 .0040 120.1	Ca3179 ppm H900.7 1.9 .2098	Mg2790 ppm H407.0 1.1 .2705
#1 #2	.0050	H1483. H1484.	0069 0017	.0313	.0005	H899.4 H902.0	H406.3 H407.8
Elem Units Avge SDev %RSD	Mn2576 ppm .9699 .0010 .1081	Co2286 ppm .0009 .0005 54.52	Fe2714 ppm .2688 .0049 1.817	Al3082 ppm 0374 .0046 12.22	V_2924 ppm 0548 .0000 .0844	Mo2020 ppm .0021 .0005 23.10	Ti3349 ppm0076 .0002 2.360
#1 #2	.9692 .9707	.0005 .0012	.2653 .2722	0341 0406	0548 0548	.0018 .0025	0075 0077
Elem Units Avge SDev %RSD	Ag3280 ppm 0005 .0003 72.31	B_2496 ppm .2649 .0011 .4120	K_7664 ppm H118.9 .4 .3455	Na3302 ppm H1608. 1.	Si2881 ppm .3922 .0010 .2577	Sn1899 ppm 0079 .0003 3.224	P_1782 ppm .1514 .0010 .6578
#1 #2	0007 0002	.2657 .2641	H119.2 H118.6	H1607. H1609.	.3930 .3915	0081 0077	.1507 .1521

Method: ZZJEM1_A Sample Name: 7985-15 x10 Operator: VVK Run Time: 05/11/11 12:52:38

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0009	.0011	.0006	.0014	.0022	0058	0052
SDev	.0022	.0058	.0001	.0007	.0001	.0004	.0051
%RSD	233.7	553.6	15.56	49.86	6.428	7.332	97.52
#1 #2	0025 .0006	.0052 0031	.0005	.0019	.0022 .0021	0061 0055	0088 0016
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0070	.0060	.0050	.0071	0230	.0017	.0054
SDev	.0088	.0001	.0004	.0025	.0042	.0002	.0060
%RSD	124.4	2.357	8.235	35.17	18.18	11.37	110.6
#1	.0132	.0061	.0053	.0054	0201	.0019	.0097
#2		.0059	.0047	.0089	0260	.0016	.0012
Elem Units Avge SDev %RSD	Sb206A ppm .0038 .0053 137.4	S_1820 ppm 92.92 .24 .2622	S_2203 ppm .0007 .0032 450.5	S_1960 ppm .0301 .0042 13.79	S_2068 ppm .0053 .0016 30.20	Ca3179 ppm H101.3 .1 .0806	Mg2790 ppm 47.05 .17
#1 #2	.0001	93.09 92.75	.0029	.0331	.0042	H101.4 H101.3	47.17 46.93
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.1238	.0003	0539	.3320	0038	.0018	.0005
SDev	.0001	.0003	.0022	.0110	.0002	.0001	.0001
%RSD	.0797	105.9	4.002	3.320	6.168	4.285	11.78
#1 #2	.1239	.0005	0523 0554	.3398 .3242	0039 0036	.0018 .0017	.0004
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0004	0094	9.276	H120.2	.1036	0001	.0456
SDev	.0004	.0015	.089	.8	.0003	.0001	.0046
%RSD	91.25	15.49	.9569	.6247	.2462	88.61	9.990
#1 #2	.0007	0105 0084	9.338 9.213	H120.8 H119.7	.1034	0001 0000	.0424

RB-01-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name:Sum	mit Environmental Tech	Contract:	Jackpile	VI.
Matrix: (soil/w	ater) <u>water</u>	Lab Sample ID:	1107985-16	
Date Received:	04/26/2011	_		
% Solids:		_		

Concentration Units (mg/L or mg/kg dry weight): mg/L

CAS No.	Analyte	Concentration	С	Q	M
7429-90-5	Aluminum	0.0105	U	100 Table 100 Ta	Р
7440-36-0	Antimony	0.0175	J		Р
7440-38-2	Arsenic	0.0033	J		P
7440-39-3	Barium	0.0009	J		P
7440-41-7	Beryllium	0.00072	Ū		Р
7440-43-9	Cadmium	0.00082	Ū		P
7440-70-2	Calcium	3.60	J	D	P
7440-47-3	Chromium	0.00109	U		P
7440-48-4	Cobalt	0.0006	J		P
7440-50-8	Copper	0.00183	U		Р
7439-89-6	Iron	0.0106	U		Р
7439-92-1	Lead	0.0027	J		Р
7439-95-4	Magnesium	0.00079	U		Р
7439-96-5	Manganese	0.00043	U		P
7440-02-0	Nickel	0.0012	J		Р
7440-09-7	Potassium	0.0240	U		P
7782-49-2	Selenium	0.0343			Р
7440-22-4	Silver	0.0029	J		Р
7440-23-5	Sodium	4.84	J	D	Р
7440-28-0	Thallium	0.00529	U		Р
7440-62-2	Vanadium	0.00174	Ū		Р
7440-66-6	Zinc	0.0058	J		P
7439-97-6	Mercury	0.00006	Ū		AV
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Com	ments:					
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Method: ZZJEM1_A Sample Name: 7985-16 Run Time: 04/29/11 19:12:18

Elem Units Avge SDev %RSD	Pb2203 ppm .0073 .0013 17.95	Pb220A ppm 0001 .0006 761.0	Cd2265 ppm .0007 .0001 18.72	Cr2677 ppm .0003 .0001 30.31	Ba4934 ppm .0009 .0000	As1890 ppm .0033 .0014 40.88	Se1960 ppm .0068 .0024 35.58
#1 #2	.0064	.0004	.0006	.0003	.0009	.0043	.0051
Elem Units Avge SDev %RSD	Se196A ppm .0073 .0018 24.07	Cu3247 ppm0012 .0008 67.88	Zn2138 ppm .0058 .0004 7.069	Ni2316 ppm .0012 .0001 7.739	Tl1908 ppm .0035 .0024 69.59	Be3130 ppm .0004 .0000 9.089	Sb2068 ppm .0167 .0083 49.60
#1 #2	.0085	0006 0018	.0061 .0055	.0012	.0018 .0052	.0004	.0226 .0108
Elem Units Avge SDev %RSD	Sb206A ppm .0164 .0086 52.62	S_1820 ppm 7352 .0470 6.399	S_2203 ppm .0027 .0000 .7268	S_1960 ppm .0343 .0004 1.066	S_2068 ppm .0175 .0085 48.65	Ca3179 ppm 1072 .0143 13.37	Mg2790 ppm 0816 .0074 9.102
#1 #2	.0225	7019 7685	.0027	.0346	.0235 .0115	0970 1173	0763 0868
Elem Units Avge SDev %RSD	Mn2576 ppm .0003 .0000 1.186	Co2286 ppm .0006 .0001 12.38	Fe2714 ppm .0038 .0003 7.643	Al3082 ppm 0390 .0018 4.512	V_2924 ppm .0006 .0005 75.24	Mo2020 ppm .0023 .0002 10.24	Ti3349 ppm .0002 .0004 176.8
#1 #2	.0003	.0005	.0036	0402 0377	.0003	.0022	.0005 0001
Elem Units Avge SDev %RSD	Ag3280 ppm .0029 .0004 12.68	B_2496 ppm .0158 .0017 10.52	K_7664 ppm 0584 .0155 26.52	Na3302 ppm .4952 .3030 61.19	Si2881 ppm .0843 .0023 2.748	Sn1899 ppm .0084 .0016 19.36	P_1782 ppm .0565 .0096 17.07
#1 #2	.0031 .0026	.0170 .0146	0475 0694	.2809 .7095	.0859 .0826	.0095	.0633

Method: ZZJEM1_A Sample Name: 7985-16 x10 Operator: VVK

Run Time: 05/11/11 12:56:12

Elem Units Avge SDev %RSD	Pb2203 ppm .0126 .0216 171.6	Pb220A ppm .0074 .0017 23.18	Cd2265 ppm 0006 .0004 67.14	Cr2677 ppm .0006 .0015 265.2	Ba4934 ppm .0005 .0000	As1890 ppm 0091 .0002 2.669	Se1960 ppm 0055 .0039 71.63
#1	.0279	.0086	0009	0005	.0005	0089	0082
#2	0027		0003	.0017	.0005	0093	0027
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm						
Avge	.0103	.0053	.0030	.0051	0369	.0021	.0098
SDev	.0016	.0001	.0001	.0007	.0051	.0001	.0038
%RSD	16.00	.8839	5.095	13.62	13.89	2.692	39.09
#1 #2	.0091 .0114	.0053 .0054	.0029	.0056	0332 0405	.0022	0125
Elem Units Avge SDev %RSD	Sb206A ppm 0072 .0002 2.143	S_1820 ppm .3116 .0013 .4270	S_2203 ppm .0094 .0083 88.63	S_1960 ppm .0339 .0048 14.19	S_2068 ppm 0005 .0012 214.0	Ca3179 ppm .3595 .0012	Mg2790 ppm 1.1269 1.0062 4.881
#1	0073	.3126	.0153	.0305	.0003	.3603	.1225
#2	0071	.3107	.0035	.0373	0014		.1313
Elem	Mn2576	Co2286 ppm0005 .0002 38.56	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm		ppm	ppm	ppm	ppm	ppm
Avge	.0013		0166	.4312	0003	0010	.0004
SDev	.0002		.0048	.0087	.0002	.0004	.0002
%RSD	17.34		29.00	2.015	63.81	37.22	51.42
#1	.0011	0004	0199	.4250	0005	0008	.0003
#2		0007	0132	.4373	0002	0013	.0006
Elem Units Avge SDev %RSD	Ag3280 ppm 0001 .0010 1592.	B_2496 ppm 0478 .0000 .0172	K_7664 ppm .1542 .0442 28.64	Na3302 (- ppm .4837 .2893 59.81	Si2881 ppm .0705 .0009 1.266	Sn1899 ppm .0001 .0006 438.8	P_1782 ppm .0203 .0066 32.37
#1	0008	0478	.1229	.2791	.0698	0003	.0156
#2	.0006	0478	.1854	.6882	.0711	.0005	.0249

RB-02-110421

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit Environmen	ntal Tech	Contract:	Jackpile
Matrix: (soil/water)wat	er	Lab Sample ID:	1107985-17
Date Received: 04/2	6/2011		-
% Solids:			
Concentration Units (mg/L or	mg/kg dry weight	:): <u>mg/L</u>	

CAS No.	Analyte	Concentration	С	Q	M
7429-90-5	Aluminum	0.0105	U	1000	P
7440-36-0	Antimony	0.0076	J		P
7440-38-2	Arsenic	0.0029	J		P
7440-39-3	Barium	0.0011	J		P
7440-41-7	Beryllium	0.00072	U	***************************************	P
7440-43-9	Cadmium	0.00082	Ū		P
7440-70-2	Calcium	1.34	J	D	Р
7440-47-3	Chromium	0.00109	U		P
7440-48-4	Cobalt	0.00051	Ū		P
7440-50-8	Copper	0.00183	U		P
7439-89-6	Iron	0.0106	Ū		Р
7439-92-1	Lead	0.00210	U		P
7439-95-4	Magnesium	0.00079	U		Р
7439-96-5	Manganese	0.00043	U		P
7440-02-0	Nickel	0.00116	U		P
7440-09-7	Potassium	0.0240	U		P
7782-49-2	Selenium	0.0257			P
7440-22-4	Silver	0.0012	J		P
7440-23-5	Sodium	2.01	J	D	P
7440-28-0	Thallium	0.0151	Ĵ		P
7440-62-2	Vanadium	0.00174	U		P
7440-66-6	Zinc	0.0067	J		P
7439-97-6	Mercury	0.00006	U		AV

Com	ments:									
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Method: ZZJEM1_A Sample Name: 7985-17 Run Time: 04/29/11 19:15:52

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm .0027 .0094 355.8	Pb220A ppm .0001 .0004 640.6	Cd2265 ppm .0002 .0001 30.09	Cr2677 ppm .0003 .0002 64.29	Ba4934 ppm .0011 .0000	As1890 ppm .0029 .0015 51.55	Se1960 ppm 0009 .0016 182.0
#1 #2	0040 .0093	0002 .0004	.0002	.0001	.0011	.0039 .0018	.0003 0020
Elem Units Avge SDev %RSD	Se196A ppm .0007 .0001 8.531	Cu3247 ppm 0017 .0011 60.61	Zn2138 ppm .0067 .0004 6.614	Ni2316 ppm 0000 .0000	Tl1908 ppm .0151 .0034 22.44	Be3130 ppm .0002 .0001 19.03	Sb2068 ppm .0084 .0034 40.67
#1 #2	.0008	0010 0025	.0070	0000 0000	.0174 .0127	.0003	.0060
Elem Units Avge SDev %RSD	Sb206A ppm .0058 .0002 2.702	S_1820 ppm 8604 .0105 1.223	S_2203 ppm .0012 .0034 283.0	S_1960 ppm .0257 .0018 7.092	S_2068 ppm .0076 .0013 16.29	Ca3179 ppm .0185 .0027 14.80	Mg2790 ppm 0952 .0014 1.451
#1 #2	.0056 .0059	8529 8678	0012 .0037	.0244	.0068	.0205 .0166	0942 0962
Elem Units Avge SDev %RSD	Mn2576 ppm .0001 .0000	Co2286 ppm .0002 .0006 251.9	Fe2714 ppm .0024 .0009 36.41	Al3082 ppm 0277 .0030 10.99	V_2924 ppm .0003 .0002 61.23	Mo2020 ppm .0016 .0001 9.366	Ti3349 ppm .0001 .0000 25.72
#1 #2	.0001	.0006 0002	.0031	0255 0298	.0004	.0017	.0001
Elem Units Avge SDev %RSD	Ag3280 ppm .0012 .0004 34.53	B_2496 ppm .0115 .0012 10.56	K_7664 ppm 0649 .0135 20.81	Na3302 ppm .1621 .0503 31.06	Si2881 ppm .0908 .0025 2.783	Sn1899 ppm .0100 .0005 5.163	P_1782 ppm .0564 .0025 4.394
#1 #2	.0014	.0106 .0124	0553 0745	.1977 .1265	.0890 .0926	.0096 .0104	.0547 .0582

Method: ZZJEM1_A Sample Name: 7985-17 x10 Run Time: 05/11/11 12:59:46 Operator: VVK

Elem Units Avge SDev %RSD	Pb2203 ppm .0059 .0048 81.93	Pb220A ppm 0037 .0001 2.318	Cd2265 ppm 0005 .0004 75.51	Cr2677 ppm .0006 .0005 88.40	Ba4934 ppm .0020 .0000	As1890 ppm 0076 .0061 79.45	Se1960 ppm .0004 .0060 1717.
#1 #2	.0025 .0093	0036 0037	0002 0008	.0002	.0020	0033 0119	.0046 0039
Elem Units Avge SDev %RSD	Se196A ppm 0055 .0026 47.00	Cu3247 ppm .0051 .0001 2.791	Zn2138 ppm 0005 .0001 20.90	Ni2316 ppm .0013 .0010 76.77	Tl1908 ppm 0043 .0074 174.1	Be3130 ppm .0023 .0002 10.57	Sb2068 ppm .0038 .0017 44.66
#1 #2	0073 0036	.0050 .0052	0004 0006	.0020 .0006	0095 .0010	.0021	.0026
Elem Units Avge SDev %RSD	Sb206A ppm 0022 .0031 141.4	S_1820 ppm .0802 .0013 1.660	S_2203 ppm 0001 .0015 1166.	S_1960 ppm .0254 .0027 10.61	S_2068 ppm .0008 .0026 332.8	Ca3179 ppm .1344 .0023 1.735	Mg2790 ppm .0312 .0000 .0016
#1 #2	0044 0000	.0792 .0811	0012 .0010	.0273 .0235	0011 .0026	.1327	.0312
Elem Units Avge SDev %RSD	Mn2576 ppm .0006 .0001 9.184	Co2286 ppm .0001 .0010 1132.	Fe2714 ppm 0179 .0108 60.43	Al3082 ppm .4512 .0081 1.798	V_2924 ppm .0004 .0000 .6884	Mo2020 ppm 0019 .0016 84.85	Ti3349 ppm .0001 .0001 94.29
#1 #2	.0006 .0006	0007 .0008	0256 0103	.4570 .4455	.0004	0030 0008	.0002
Elem Units Avge SDev %RSD	Ag3280 ppm .0002 .0000 12.48	B_2496 ppm 0505 .0002 .3261	K_7664 ppm .1086 .0064 5.928	Na3302 ppm .2005 .0881 43.96	Si2881 ppm .0696 .0009 1.282	Sn1899 ppm 0004 .0006 169.7	P_1782 ppm .0198 .0013 6.723
#1 #2	.0002	0504 0506	.1041	.1382 .2628	.0702 .0689	.0001	.0188 .0207

RM-JM-SED-02-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit En	vironmental Tech	Contract:	Jackpile	
Matrix: (soil/water)	soil	_ Lab Sample ID: _	1107985-18	
Date Received:	04/26/2011	_		
Solids:	100	_		

Concentration Units (mg/L or mg/kg dry weight): mg/kg

CAS No.	Analyte	Concentration	C	Q	М
7429-90-5	Aluminum	7420		D	Р
7440-36-0	Antimony	0.529	J		Р
7440-38-2	Arsenic	4.65			Р
7440-39-3	Barium	68.0			P
7440-41-7	Beryllium	0.393	J		Р
7440-43-9	Cadmium	0.240	J		Р
7440-70-2	Calcium	20900		D	P
7440-47-3	Chromium	7.88			Р
7440-48-4	Cobalt	5.22			P
7440-50-8	Copper	8.27			P
7439-89-6	Iron	17900		D	₽
7439-92-1	Lead	3.06	J		P
7439-95-4	Magnesium	5060			P
7439-96-5	Manganese	160			P
7440-02-0	Nickel	9.36			P
7440-09-7	Potassium	2310			P
7782-49-2	Selenium	0.109	Ū		P
7440-22-4	Silver	0.109	Ū		Р
7440-23-5	Sodium	3880			Р
7440-28-0	Thallium	0.174	Ū		Р
7440-62-2	Vanadium	19.8			Р
7440-66-6	Zinc	34.2			Р
7439-97-6	Mercury	0.0436	U		AV

Comr	ments:									
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Method: ZZJEM1_A Sample Name: 7985-18s

Run Time: 05/12/11 13:05:36

Elem	Pb2203	Pb220A	Cd2265 ppm .0044 .0003 7.338	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm		ppm	ppm	ppm	ppm
Avge	.1028	.1013		.1445	1.247	.0852	.5487
SDev	.0008	.0040		.0004	.003	.0049	.0040
%RSD	.7718	3.955		.2600	.2740	5.767	.7309
#1 #2	.1034	.0984 .1041	.0042	.1448	1.244	.0887	.5515 .5458
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908 ppm0500 .0250 50.07	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm		ppm	ppm
Avge	<-1.000	.1517	.6266	.1716		.0072	.0141
SDev	.002	.0001	.0022	.0020		.0001	.0085
%RSD	.0989	.0679	.3593	1.154		.8536	60.40
#1 #2	<-1.000 <-1.000	.1517	.6250 .6282	.1730 .1702	0677 0323	.0072	.0202
Elem Units Avge SDev %RSD	Sb206A ppm .0056 .0042 75.61	S_1820 ppm 66.20 .22 .3326	S_2203 ppm .0562 .0023 4.109	S_1960 ppm 7321 .0012 .1606	S_2068 ppm .0097 .0056 57.60	Ca3179 ppm 332.3 1.4	Mg2790 ppm 92.77 √ .36 .3863
#1	.0086	66.04	.0546	7329	.0136	331.3	92.51
#2	.0026	66.35	.0578	7313	.0057	333.3	93.02
Elem Units Avge SDev %RSD	Mn2576 ppm 2.937 .011 .3796	Co2286 ppm .0957 .0005 .5071	Fe2714 ppm 288.5 .9	A13082 / ppm 135.6 / .4 / .3163	V_2924 ppm .3626 \ .0016 .4394	Mo2020 ppm .0059 .0016 27.05	Ti3349 ppm 4.739 .015 .3270
#1	2.930	.0953	287.9	135.3	.3615	.0048	4.729
#2	2.945	.0960	289.1	135.9	.3637		4.750
Elem Units Avge SDev %RSD	Ag3280 ppm 0022 .0009 V	B_2496 ppm .5940 .0006 .0929	K_7664 ppm 42.43 .14	Na3302 ppm 71.23 .12 .1720	Si2881 ppm 5.276 .019 .3527	Sn1899 ppm .0105 .0008 7.458	P_1782 ppm 4.482 .018 .3938
#1	0016	.5944	42.33	71.14	5.263	.0099	4.469
#2	0028	.5936	42.53	71.31	5.289		4.494

Method: ZZJEM1_A Sample Name: 7985-18s x10 Operator: VVK

Run Time: 05/12/11 13:10:33

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0030	.0103	0015	.0148	.1177	.0149	.0678
SDev	.0118	.0010	.0003	.0008	.0008	.0017	.0017
%RSD	397.5	9.376	21.76	5.692	.6449	11.56	2.514
#1	0113	.0109	0013	.0154	.1182	.0136	.0690
#2	.0054	.0096	0018		.1172	.0161	.0666
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	2230	.0162	.0741	.0186	.0096	.0012	.0018
SDev	.0097	.0001	.0009	.0006	.0004	.0001	.0033
%RSD	4.352	.9513	1.275	3.276	4.562	8.111	184.6
#1 #2	2162 2299	.0161 .0164	.0748 .0735	.0190 .0182	.0099	.0011	0005 .0041
Elem Units Avge SDev %RSD	Sb206A ppm 0013 .0012 88.39	S_1820 ppm 8.350 .042 .5000	S_2203 ppm .0009 .0033 366.8	S_1960 ppm 0533 .0073 13.78	S_2068 ppm .0008 .0002 27.37	Ca3179 ppm 38.26 .30 .7878	Mg2790 ppm 10.30 .06
#1	0005	8.380	0014	0481	.0006	38.48	10.34
#2	0022	8.321	.0032	0585		38.05	10.25
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.3334	.0093	32.91	13.61	.0375	0001	.5132
SDev	.0019	.0000	.22	.07	.0000	.0028	.0037
%RSD	.5720	.0025	.6647	.4853	.0146	2758.	.7228
#1	.3347	.0093	33.06	13.66	.0375	.0019	.5158
#2	.3320	.0093	32.75	13.57		0021	.5106
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0015	.0702	3.049	5.710	.6284	.0011	.5526
SDev	.0000	.0009	.024	.162	.0058	.0002	.0002
%RSD	1.433	1.270	.7937	2.834	.9175	20.50	.0303
#1	0016	.0708	3.066	5.825	.6324	.0013	.5527
#2	0015	.0696	3.032	5.596	.6243		.5524

RM-JM-SED-110420

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1A-IN INORGANIC ANALYSIS DATA SHEET

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Lab Name: _	Summit Envir	conmental Te	ech Contr	ract: _		Jackpi.	le
Matrix: (soi	il/water)	soil	Lab S	Sample	ID:	110798	35-19
Date Receive	ed:	04/26/2011					
	8						
Concentratio	on Units (mg/	L or mg/kg	dry weight): _n	ng/kg			
	CAS No.	Analyte	Concentration	С	Q)	М
	7429-90-5	Aluminum	6660		D		Р
	7440-36-0	Antimony	23.3		N		P
	7440-38-2	Arsenic	5.72				P
	7440-39-3	Barium	57.0				Р
	7440-41-7	Beryllium	0.558	J	*******	***************************************	P
	7440-43-9	Cadmium	0.0237	U			P
	7440-70-2	Calcium	18000		D		P
	7440-47-3	Chromium	9.10			······································	P
	7440-48-4	Cobalt	5.26				P
	7440-50-8	Copper	7.31				Р
	7439-89-6	Iron	16900		D		P
	7439-92-1	Lead	2.35	J			P
	7439-95-4	Magnesium	4890				P
	7439-96-5	Manganese	173				P
	7440-02-0	Nickel	9.18		N		P
	7440-09-7	Potassium	2660		N		P
	7782-49-2	Selenium	23.3				P
	7440-22-4	Silver	0.119	U :	N,*		P
	7440-23-5	Sodium	4240		D		P

0.717

0.0474

22.4

36

7440-28-0 Thallium

7440-62-2 Vanadium

7439-97-6 Mercury

7440-66-6 Zinc

Comments:

59.31

Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-19s

Run Time: 04/29/11 19:55:08

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Mode: CONC Corr. Factor: 1

				_	p == 1		U
Elem Units Avge SDev %RSD	Pb2203 ppm .1586 .0050 3.153	Pb220A ppm .0483 .0010 2.155	Cd2265 ppm 0235 .0002 .6575	Cr2677 ppm .1535 .0001 / \(^0\)	Ba4934 ppm 0 .9614 1 .0011 1	As1890 ppm .0964 .0027 2.842	Se1960 ppm .3315 .0031 .9258
#1 #2	.1550 .1621	.0491 .0476	0237 0234	.1534 .1535	.9621 .9606	.0983 .0945	.3293 .3336
Elem Units Avge SDev %RSD	Se196A ppm 2136 .0136 6.372	Cu3247 ppm .123313 .0015 1.257	Zn2138 ppm .6068 24 .0013	Ni2316 ppm .1548 .0001 .0375	T11908 ppm .0121 .0002	Be3130 ppm .0094 .0000 .2557	Sb2068 ppm .0136 .0086 63.55
#1 #2	2040 2232	.1244	.6078 .6059	.1548	.0122 .0119	.0095 .0094	.0197
Elem Units Avge SDev %RSD	Sb206A ppm 0443 .0039 8.897	S_1820 ppm H151.9 .5 .3237	S_2203 ppm .0397 .0009 2.405	S_1960 ppm .3935 .0087 2.205	S_2068 ppm	Ca3179 ppm H359.8 .6	Mg2790 ppm 82.38 .16 .1968
#1 #2		H152.3 H151.6	.0390	.3997 .3874		H360.3 H359.4	82.50 82.27
Elem Units Avge SDev %RSD	Mn2576 ppm 2.916 .005 .1726	Co2286, 7 ppm 7 .0887 .0002 .2251	Fe2714 ppm H287.6 .5 .1573	A13082 ppm H130.2 .3	V_2924 ppm .3775 .0007 .1807	Mo2020 ppm .0165 .0023 13.68	Ti3349 ppm 3.656 .007 .2043
#1 #2	2.920 2.913		H287.9 H287.2	H130.4 H130.0 $\sqrt{}$.3780 .3771	.0181 .0149	3.662 3.651
Elem Units Avge SDev %RSD	Ag3280 ppm .0001 .0001	B_2496 ppm .5719 .0007 .1191	K_7664 ppm 44.907 .03 .0654	Na3302 ppm H175.1 .4	Si2881 ppm 5.206 .010 .1990	Sn1899 ppm .0408 .0019 4.785	P_1782 ppm 7.464 .003 .0443
#1 #2	.0000	.5723 .5714	44.88 44.92	H175.4 H174.8	5.214 5.199	.0422	7.466 7.461

refret + some mg/l × 0.000 l = 50

Method: ZZJEM1_A Sample Name: 7985-19s x10 Operator: VVK Run Time: 05/11/11 14:45:43

Comment: 6010/200.7 CPI 4400-013 lot 10B101 (2)
Mode: CONC Corr. Factor: 1

59	3	
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Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0053	.0209	.0008	.0166	.0880	.0047	.0131
SDev	.0011	.0056	.0002	.0064	.0373	.0040	.0107
%RSD	21.66	26.97	24.70	38.28	42.35	84.91	81.27
#1 #2	.0061 .0045	.0249	.0009	.0211	.1144 .0617	.0076 .0019	.0056 .0207
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	1559	.0158	.0644	.0189	0254	.0027	.0165
SDev	.0599	.0053	.0293	.0075	.0124	.0001	.0095
%RSD	38.42	33.36	45.48	39.73	48.66	4.215	57.77
#1 #2	1983 1136	.0195 .0120	.0852 .0437	.0243	0167 0342	.0027 .0026	.0233
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0014	8.127	.0114	0328	.0056	30.43	8.075
SDev	.0008	3.422	.0022	.0261	.0027	13.43	3.543
%RSD	56.57	42.11	18.75	79.68	47.58	44.12	43.87
#1	0020	10.55	.0130	0513	.0075	39.92 ·	10.58
#2	0008	5.707	.0099	0143	.0037	20.94	5.570
Elem	Mn2576	Co2286 ppm .0105 .0045 42.75	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm		ppm	ppm	7ppm	ppm	ppm
Avge	.2883		28.53	11.23	.0374	.0037	.3436
SDev	.1246		12.53	4.81	.0159	.0014	.1493
%RSD	43.22		43.91	42.86	42.60	36.55	43.45
#1 #2	.3764	.0137	37.39 19.67	14.63 7.828	.0487 .0261	.0047 .0028	.4492 .2381
Elem	Ag3280	B_2496	K_7664	Na3302 (ppm 7.153 () 3.489 48.77	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm) ppm	ppm	ppm
Avge	.0021	.0406	2.645		.5516	.0015	.5434
SDev	.0012	.0295	1.231		.2241	.0004	.2464
%RSD	57.71	72.67	46.55		40.63	23.79	45.34
#1 #2	.0030	.0614 .0197	3.516 1.775	9.620 4.687	.7100 .3931	.0018	.7176 .3692

RM-JM-SW-02-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

ORGANIC	ANALYSIS	DATA	SHEET	

Lab Name:	Summit Envir	onmental Te	ch Contr	act:	Jack	pile
Matrix: (soi	l/water)	water	Lab S	ample	- TD· 110	7985-20
				op _ c		7505-20
Date Receive	d:	04/26/2011				
% Solids:						

Concentration	n Units (mg/	L or mg/kg	dry weight): _m	g/L	_	
	CAS No.	Analyte	Concentration	С	Q	M
	7429-90-5	Aluminum	0.0105	U	10000	P
	7440-36-0	Antimony	0.0062	J		P
	7440-38-2	Arsenic	0.00206	Ū		P
	7440-39-3	Barium	0.0232	J		P
	7440-41-7	Beryllium	0.00072	Ū		P
	7440-43-9	Cadmium	0.00082	U		P
	7440-70-2	Calcium	224		D	P
	7440-47-3	Chromium	0.00109	U		P
	7440-48-4	Cobalt	0.00051	U		P
	7440-50-8	Copper	0.00183	U		P
	7439-89-6	Iron	0.0106	Ū		P
	7439-92-1	Lead	0.00210	Ū		P
	7439-95-4	Magnesium	222		D	P
	7439-96-5	Manganese	0.0152	J		P
	7440-02-0	Nickel	0.0023	J		P
	7440-09-7	Potassium	18.6			P
	7782-49-2	Selenium	0.0278			P
	7440-22-4	Silver	0.00039	U		P
	7440-23-5	Sodium	532		D	P
	7440-28-0	Thallium	0.0154	J		P
	7440-62-2	Vanadium	0.00174	Ū		Р
	7440-66-6	Zinc	0.00053	Ų		Р
	7439-97-6	Mercury	0.00006	Ū		AV

Comments	:					
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Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-20 Run Time: 04/29/11 19:19:27

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm .0003 .0009 339.7	Pb220A ppm 0018 .0016 89.59	Cd2265 ppm .0003 .0002 77.28	Cr2677 ppm 0002 .0003 125.8	Ba4934 ppm .0232 .0001 .4637	As1890 ppm .0008 .0040 528.0	Se1960 ppm .0090 .0033 36.31
#1 #2	0004 .0009	0029 0007	.0004	0004 0000	.0231	0021 .0036	.0114
Elem Units Avge SDev %RSD	Se196A ppm 0037 .0002 6.337	Cu3247 ppm 0037 .0007 20.20	Zn2138 ppm 0014 .0000 2.190	Ni2316 ppm .0023 .0006 26.40	Tl1908 ppm .0154 .0039 25.02	Be3130 ppm .0001 .0000 35.23	Sb2068 ppm .0022 .0091 405.3
#1 #2	0035 0038	0031 0042	0014 0014	.0019	.0127 .0181	.0001	0042 .0087
Elem Units Avge SDev %RSD	Sb206A ppm .0067 .0069 102.1	S_1820 ppm H596.6 .9 .1472	S_2203 ppm 0008 .0014 182.2	S_1960 ppm .0278 .0013 4.491	S_2068 ppm .0062 .0015 23.64	Ca3179 ppm H225.8 .5 .2146	Mg2790 ppm H206.8 .5 .2223
#1 #2	.0116	H596.0 H597.2	0018 .0002	.0287 .0269	.0072 .0051	H225.4 H226.1	H206.5 H207.1
Elem Units Avge SDev %RSD	Mn2576 ppm .0152 .0001 .3525	Co2286 ppm .0001 .0001 78.98	Fe2714 ppm 0106 .0008 7.834	Al3082 ppm 0393 .0058 14.86	V_2924 ppm 0271 .0002 .5906	Mo2020 ppm .0043 .0005 11.49	Ti3349 ppm 0017 .0000 1.757
#1 #2	.0152 .0151	.0001	0112 0100	0435 0352	0269 0272	.0039	0017 0017
Elem Units Avge SDev %RSD	Ag3280 ppm .0002 .0001 99.97	B_2496 ppm .1919 .0015 .8100	K_7664 ppm 18.58 .03	Na3302 ppm H760.2 1.6 .2042	Si2881 ppm 4.658 .002 .0479	Sn1899 ppm .0028 .0012 42.35	P_1782 ppm .0662 .0010 1.447
#1 #2	.0000	.1930 .1908	18.60 18.57	H759.1 H761.3	4.656 4.660	.0036	.0669 .0655

Method: ZZJEM1_A Sample Name: 7985-20 x10 Run Time: 05/11/11 13:03:20 Comment: 6010/200.7 CPI 4400-013 lot 10B101 Operator: VVK

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0056	.0027	0003	.0007	.0029	0093	.0047
SDev	.0050	.0024	.0001	.0007	.0001	.0044	.0061
%RSD	89.16	88.58	48.37	99.03	2.397	46.71	128.5
#1 #2	.0092 .0021	.0010	0002 0004	.0012 .0002	.0028 .0029	0124 0063	.0090
Elem	Se196A	Cu3247	Zn2138	Ni2316	T11908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0035	.0053	.0005	.0057	.0079	.0023	.0033
SDev	.0109	.0001	.0008	.0016	.0200	.0000	.0016
%RSD	309.1	.8951	167.2	28.28	253.4	.2933	48.55
#1 #2	0042 .0112	.0052	0001 .0011	.0045	0063 .0221	.0023	.0044
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0005	32.11	.0040	.0345	.0025	22.44	22.24
SDev	.0010	.05	.0000	.0100	.0011	.02	.03
%RSD	204.3	.1483	.8600	29.14	44.88	.0727	.1353
#1	.0012	32.14	.0040	.0274	.0033	22.43	22.26
#2	0002	32.07		.0416	.0017	22.45	22.22
Elem Units Avge SDev %RSD	Mn2576 ppm .0031 .0001 1.774	Co2286 ppm0000 .0002 420.8	Fe2714 ppm0554 .0018 3.266	Al3082 ppm .4598 .0017 .3764	V_2924 ppm 0015 .0002 13.97	Mo2020 ppm 0013 .0012 94.28	Ti3349 ppm0000 .0003 4401000.
#1	.0031	0002	0541	.4611	0014	0004	0002
#2	.0032	.0001	0566	.4586	0017	0022	.0002
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0001	0301	1.537	53.19	.6072	0001	.0190
SDev	.0001	.0004	.042	.09	.0004	.0011	.0023
%RSD	130.9	1.203	2.754	.1693	.0630	1020.	12.24
#1	.0000	0304	1.507	53.12	.6075	0009	.0207
#2		0298	1.566	53.25	.6070	.0007	.0174

RM-JM-SW-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name:	Summit Envir	conmental Te	ch Contr	act:	Jackpi	.le			
Matrix: (soil Date Received	· —	water 04/26/2011		ample	ID:11079	85-21			
% Solids:									
Concentration	u Units (mg/	L or mg/kg	dry weight): _m	g/L	-				
	CAS No.	Analyte	Concentration	С	Q	M			
	7429-90-5 Aluminum 0.0105 U P								
	7440-36-0	Antimony	0.0041	J		Р			
	7440-38-2	Arsenic	0.00206	Ū		Р			
	7440-39-3	Barium	0 0228	I.T		D			

7440-36-0	Antimony	0.0041	J		P
7440-38-2	Arsenic	0.00206	U		Р
7440-39-3	Barium	0.0228	J		P
7440-41-7	Beryllium	0.00072	U		Р
7440-43-9	Cadmium	0.00082	U		Р
7440-70-2	Calcium	218		D	P
7440-47-3	Chromium	0.00109	U		Р
7440-48-4	Cobalt	0.00051	Ū		P
7440-50-8	Copper	0.00183	U	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Р
7439-89-6	Iron	0.0106	U		P
7439-92-1	Lead	0.0025	J		Р
7439-95-4	Magnesium	217		D	P
7439-96-5	Manganese	0.0131	J		Р
7440-02-0	Nickel	0.00116	Ū		P
7440-09-7	Potassium	18.1			P
7782-49-2	Selenium	0.0324			P
7440-22-4	Silver	0.00039	U		P
7440-23-5	Sodium	519		D	P
7440-28-0	Thallium	0.0175	J		Р
7440-62-2	Vanadium	0.00174	Ū		Р
7440-66-6	Zinc	0.00053	U		Р
7439-97-6	Mercury	0.00006	U		AV

Com	ments:							
		 	 		 	—- <u>-</u>	 	
,		 	 				 	
		 	 	1010				

Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-21 Run Time: 04/29/11 19:23:01

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0121 .0083 69.01	Pb220A ppm 0027 .0045 165.0	Cd2265 ppm .0002 .0001 65.67	Cr2677 ppm 0002 .0004 240.8	Ba4934 ppm .0228 .0002 .9460	As1890 ppm .0007 .0010 157.1	Se1960 ppm .0049 .0010 21.29
#1 #2	.0180 .0062	.0005 0059	.0003	.0001	.0229	.0014	.0056 .0041
Elem Units Avge SDev %RSD	Se196A ppm .0054 .0030 55.06	Cu3247 ppm0032 .0007 20.83	Zn2138 ppm 0022 .0002 7.623	Ni2316 ppm .0014 .0008 53.85	Tl1908 ppm .0175 .0073 41.75	Be3130 ppm 0000 .0000 127.6	Sb2068 ppm 0028 .0050 179.0
#1 #2	.0033	0027 0037	0021 0023	.0019	.0123 .0227	0000 0001	.0007 0063
Elem Units Avge SDev %RSD	Sb206A ppm .0063 .0056 88.13	S_1820 ppm H582.3 2.4 .4205	S_2203 ppm .0025 .0058 229.9	S_1960 ppm .0324 .0016 5.024	S_2068 ppm .0041 .0019 45.07	Ca3179 ppm H220.7 .8	Mg2790 ppm H202.3 .9
#1 #2	.0024	H580.6 H584.0	.0066 0016	.0313	.0028 .0055	H220.1 H221.2	H201.7 H202.9
Elem Units Avge SDev %RSD	Mn2576 ppm .0131 .0001 .7751	Co2286 ppm .0003 .0006 184.1	Fe2714 ppm 0075 .0041 54.42	Al3082 ppm 0412 .0018 4.271	V_2924 ppm 0258 .0005 1.826	Mo2020 ppm .0050 .0004 7.741	Ti3349 ppm0019 .0004 23.44
#1 #2	.0132 .0130	.0007 0001	0046 0103	0399 0424	0255 0262	.0053	0016 0022
Elem Units Avge SDev %RSD	Ag3280 ppm 0003 .0007 210.9	B_2496 ppm .1870 .0034 1.834	K_7664 ppm 18.07 .08 .4546	Na3302 ppm H743.5 2.9 .3840	Si2881 ppm 4.541 .017 .3830	Sn1899 ppm .0019 .0012 64.96	P_1782 ppm .0622 .0000
#1 #2	.0002	.1895 .1846	18.02 18.13	H741.5 H745.5	4.529 4.554	.0010	.0622 .0622

Method: ZZJEM1_A Sample Name: 7985-21 x10 Run Time: 05/11/11 13:06:54 Comment: 6010/200.7 CPI 4400-013 lot 10B101 Operator: VVK

Elem Units Avge SDev %RSD	Pb2203 ppm 0042 .0062 148.3	Pb220A ppm .0016 .0052 331.8	Cd2265 ppm .0003 .0006 199.3	Cr2677 ppm .0011 .0000 .0001	Ba4934 ppm .0028 .0001 2.481	As1890 ppm 0072 .0026 36.42	Se1960 ppm 0016 .0032 199.5
#1 #2	.0002	.0052 0021	0001 .0007	.0011	.0028 .0027	0090 0053	.0007 0039
Elem Units Avge SDev %RSD	Se196A ppm .0059 .0019 33.03	Cu3247 ppm .0057 .0002 3.308	Zn2138 ppm .0026 .0001 1.750	Ni2316 ppm .0090 .0020 22.58	Tl1908 ppm 0161 .0400 248.2	Be3130 ppm .0024 .0000	Sb2068 ppm .0028 .0060 217.2
#1 #2	.0045	.0056 .0058	.0026 .0025	.0076 .0104	.0122	.0023	.0070 0015
Elem Units Avge SDev %RSD	Sb206A ppm 0038 .0037 96.97	S_1820 ppm 31.33 .04 .1310	S_2203 ppm 0001 .0056 11030.	S_1960 ppm .0322 .0022 6.855	S_2068 ppm 0006 .0005 75.16	Ca3179 ppm 21.77 .02 .0857	Mg2790 ppm 21.71 .04 .1752
#1 #2	0065 0012	31.36 31.30	.0039	.0338	0010 0003	21.79 21.76	21.73 21.68
Elem Units Avge SDev %RSD	Mn2576 ppm .0030 .0000	Co2286 ppm0003 .0005 188.3	Fe2714 ppm 0585 .0150 25.57	Al3082 ppm .4754 .0052 1.090	V_2924 ppm 0016 .0008 50.72	Mo2020 ppm 0007 .0001 10.88	Ti3349 ppm 0000 .0001 283.0
#1 #2	.0029	0007 .0001	0479 0690	.4791 .4717	0010 0022	0008 0007	.0000 0001
Elem Units Avge SDev %RSD	Ag3280 ppm .0000 .0000	B_2496 ppm 0307 .0009 3.058	K_7664 ppm 1.566 .001 .0886	Na3302 ppm 51.90 .39 .7493	Si2881 ppm .5964 .0014 .2351	Sn1899 ppm .0009 .0008 89.96	P_1782 ppm .0182 .0063 34.70
#1 #2	.0000	0301 0314	1.565 1.567	52.18 51.63	.5954 .5974	.0003	.0137

RM-SED-BG-110419

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit Env	ronmental Tech	Contract:	Jackpile	
Matrix: (soil/water)	soil	Lab Sample ID: _	1107985-22	
Date Received:	04/26/2011	_		
% Solids:	70.99	_		

Concentration Units (mg/L or mg/kg dry weight): mg/kg

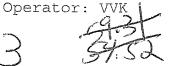
CAS No.	Analyte	Concentration	C	Q	М
7429-90-5	Aluminum	15800		D	ΙP
7440-36-0	Antimony	0.113	Ū		P
7440-38-2	Arsenic	6.68			P
7440-39-3	Barium	154			P
7440-41-7	Beryllium	0.916	J		P
7440-43-9	Cadmium	0.0282	Ū		P
7440-70-2	Calcium	23300		D	Р
7440-47-3	Chromium	8.40			P
7440-48-4	Cobalt	7.78			P
7440-50-8	Copper	13.1			P
7439-89-6	Iron	25900		D	P
7439-92-1	Lead	4.58	J		P
7439-95-4	Magnesium	6910			P
7439-96-5	Manganese	226			P
7440-02-0	Nickel	14.6			P
7440-09-7	Potassium	4180			P
7782-49-2	Selenium	29.1			P
7440-22-4	Silver	0.141	Ū		P
7440-23-5	Sodium	1950		D	P
7440-28-0	Thallium	0.662	J		P
7440-62-2	Vanadium	29.8			P
7440-66-6	Zinc	57.0			P
7439-97-6	Mercury	0.0563	Ū		AV

Comments:			
		77.50	
	74 Add		

Method: ZZJEM1_A Sample Name: 7985-22s

Run Time: 04/29/11 20:05:51 Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1





Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm , 0	ppm	ppm	ppm	ppm
Avge	.2295	.0531	0235	.1992	2.189	.0948	.3414
SDev	.0008	.0000	.0001	.0001	.002	.0041	.0034
%RSD	.3510	.0758	.5129	.0667	.1106	4.300	.9822
#1	.2301	.0532	0236	.1993	2.191	.0919	.3391
#2	.2290	.0531	0235	.1991	2.188	.0977	.3438
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	2087	.1858	.8098	.2080	.0094	.0130	0023
SDev	.0070	.0004	.0024	.0006	.0159	.0000	.0024
%RSD	3.335	.2086	.2993	.2635	168.0	.2766	105.6
#1	2137	.1860	.8115	.2076	.0207	.0130	0006
#2	2038	.1855	.8081	.2084	0018	.0129	0040
Elem Units Avge SDev %RSD	Sb206A ppm 0458 .0024 5.204	S_1820 ppm 19.60 .08 .4224	S_2203 ppm .0650 .0002 .3669	S_1960 ppm .4136 .0051 1.225	S_2068 ppm 0301 .0025 8.290	Ca3179 ppm H311.7 .5	Mg2790 ppm 98.13 .16
#1	0441	19.66	.0652	.4101	0283	H312.1	98.24
#2	0475	19.54	.0648		0318	H311.4	98.01
Elem Units Avge SDev %RSD	Mn2576 ppm 3.215 .005 .1551	Co2286 ppm .1104 .0008 .7574	Fe2714 ppm H297.3 .5	Al3082 ppm H203.8 .4 .1794	V_2924 ppm .4234 .0007 .1758	Mo2020 ppm .0154 .0004 2.520	Ti3349 ppm 1.344 .003 .2367
#1	3.219	.1109	H297.7	H204.1	.4229	.0156	1.346
#2	3.212	.1098	H297.0	H203.5	.4239	.0151	1.342
Elem Units Avge SDev %RSD	Ag3280 ppm 0020 .0004 19.87	B_2496 ppm .5990 .0032 .5338	K_7664 ppm 59.37 .10	Na3302 ppm 39.20 .17 .4316	Si2881 ppm 4.626 .016 .3478	Sn1899 ppm .0483 .0008 1.599	P_1782 ppm 8.034 .004 .0524
#1	0022	.6013	59.44	39.08	4.637	.0488	8.031
#2	0017	.5967	59.31	39.32	4.614	.0477	8.037

Method: ZZJEM1_A Sample Name: 7985-22s x10 Operator: VVK Run Time: 05/11/11 14:49:17 Comment: 6010/200.7 CPI 4400-013 lot 10B101

70	λ_j	9

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0223	.0211	.0001	.0251	.2512	.0067	.0367
SDev	.0105	.0005	.0001	.0002	.0017	.0021	.0181
%RSD	47.27	2.481	90.63	.8950	.6871	31.88	49.34
#1 #2	.0148	.0208 .0215	.0000	.0253 .0250	.2524 .2500	.0052 .0082	.0495
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	2248	.0252	.1578	.0334	0161	.0032	.0084
SDev	.0049	.0003	.0037	.0042	.0015	.0000	.0076
%RSD	2.175	1.247	2.369	12.48	9.588	1.124	91.01
#1 #2	2283 2213	.0255	.1552 .1605	.0304	0172 0150	.0032 .0032	.0137
Elem Units Avge SDev %RSD	Sb206A ppm .0008 .0046 574.0	S_1820 ppm 1.465 .010 .7007	S_2203 ppm .0161 .0038 23.83	S_1960 ppm 0578 .0056 9.755	S_2068 ppm .0043 .0056 128.6	Ca3179 ppm 33.02 .25 .7442	Mg2790 ppm 11.84 .11
#1 #2	.0040	1.472	.0134	0538 0617	.0083	33.19 32.84	11.92 11.77
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349 ppm .1588 .0015 .9165
Units	ppm	ppm	ppm	ppm	ppm	ppm	
Avge	.3974	.0145	36.79	22.41	.0510	.0038	
SDev	.0036	.0001	.33	.17	.0000	.0018	
%RSD	.9157	.6885	.8909	.7771	.0014	47.14	
#1	.4000	.0144	37.02	22.53	.0510	.0025	.1598
#2	.3948	.0145	36.56	22.29	.0510	.0050	.1577
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0002	.0498	5.616	2.765	.6155	.0022	.8077
SDev	.0002	.0019	.036	.021	.0014	.0003	.0031
%RSD	97.33	3.869	.6491	.7738	.2230	11.25	.3823
#1 #2	.0001	.0511	5.590 5.641	2.780 2.750	.6164 .6145	.0024	.8055 .8099

RM-SW-BG-110419

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit Environmental	Tech Contract:	Jackpile
Matrix: (soil/water)water	Lab Sample :	ID:1107985-23
Date Received: 04/26/20	11	
% Solids:		
Concentration Units (mg/L or mg/l	g dry weight): mg/L	

CAS No.	Analyte	Concentration	С	Q	M
7429-90-5	Aluminum	0.0105	U		Р
7440-36-0	Antimony	0.0043	J		P
7440-38-2	Arsenic	0.00206	Ü		Р
7440-39-3	Barium	0.0261	J		P
7440-41-7	Beryllium	0.00072	U		Р
7440-43-9	Cadmium	0.00082	U		P
7440-70-2	Calcium	117		D	Р
7440-47-3	Chromium	0.00109	U		P
7440-48-4	Cobalt	0.00051	U		Р
7440-50-8	Copper	0.00183	U		P
7439-89-6	Iron	0.0106	U		P
7439-92-1	Lead	0.00210	Ū		Р
7439-95-4	Magnesium	106		D	P
7439-96-5	Manganese	0.00043	Ū		P
7440-02-0	Nickel	0.00116	U		P
7440-09-7	Potassium	15.0			P
7782-49-2	Selenium	0.0350			P
7440-22-4	Silver	0.00039	Ū		P
7440-23-5	Sodium	280		D	P
7440-28-0	Thallium	0.0151	J		P
7440-62-2	Vanadium	0.001741	U		P
7440-66-6	Zinc	0.0023	J		P
7439-97-6	Mercury	0.00006	U		AV

Comn	ments:				
_			VIII) _ VIII		
_		 		 	

Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-23 Run Time: 04/29/11 19:26:35

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm . .0092 .0040 42.77	Pb220A ppm 0033 .0008 23.05	Cd2265 ppm .0002 .0002 140.4	Cr2677 ppm 0001 .0000 47.59	Ba4934 ppm .0261 .0000	As1890 ppm 0019 .0002 12.99	Se1960 ppm .0155 .0034 21.64
#1 #2	.0120 .0065	0039 0028	.0003	0000 0001	.0261 .0261	0017 0020	.0131 .0179
Elem Units Avge SDev %RSD	Se196A ppm .0040 .0028 70.21	Cu3247 ppm 0037 .0004 11.47	Zn2138 ppm .0023 .0002 10.09	Ni2316 ppm .0003 .0004 152.7	Tl1908 ppm .0151 .0159 105.0	Be3130 ppm 0000 .0000 458.1	Sb2068 ppm0042 .0003 6.404
#1 #2	.0060 .0020	0040 0034	.0025 .0021	.0006 0000	.0263	.0000	0040 0044
Elem Units Avge SDev %RSD	Sb206A ppm .0071 .0011 15.06	S_1820 ppm H300.2 2.2 .7407	S_2203 ppm .0012 .0008 67.14	S_1960 ppm .0350 .0008 2.170	S_2068 ppm .0043 .0006 14.40	Ca3179 ppm H121.3 .8	Mg2790 ppm H100.9 .9
#1 #2	.0063 .0078	H298.6 H301.8	.0017	.0355	.0039	H120.7 H121.9	H100.3 H101.6
Elem Units Avge SDev %RSD	Mn2576 ppm0079 .0001 1.045	Co2286 ppm 0000 .0005 1793.	Fe2714 ppm0587 .0025 4.334	Al3082 ppm 0262 .0038 14.54	V_2924 ppm 0145 .0000 .0556	Mo2020 ppm .0035 .0006 17.68	Ti3349 ppm0012 .0001 7.253
#1 #2	0078 0080	.0003 0004	0569 0605	0289 0235	0145 0145	.0039	0012 0013
Elem Units Avge SDev %RSD	Ag3280 ppm .0002 .0006 261.6	B_2496 ppm .1400 .0010 .7050	K_7664 ppm 15.02 .07 .4808	Na3302 ppm H406.9 3.4 .8353	Si2881 ppm 4.473 .033 .7382	Sn1899 ppm .0067 .0026 38.91	P_1782 ppm .0726 .0019 2.667
#1 #2	0002 .0007	.1393 .1407	14.97 15.08	H404.5 H409.3	4.449 4.496	.0085	.0712

Method: ZZJEM1_A Sample Name: 7985-23 x10 Run Time: 05/11/11 13:10:28 Comment: 6010/200.7 CPI 4400-013 lot 10B101 Operator: VVK

Elem Units Avge SDev %RSD	Pb2203 ppm 0040 .0046 116.6	Pb220A ppm .0011 .0053 470.1	Cd2265 ppm 0004 .0008 188.1	Cr2677 ppm .0004 .0002 56.57	Ba4934 ppm .0029 .0000	As1890 ppm 0071 .0032 45.43	Se1960 ppm .0026 .0091 348.6
#1 #2	0072 0007	0026 .0048	0009 .0001	.0005	.0029 .0029	0048 0093	0038 .0090
Elem Unit: Avge SDev %RSD	s ppm .0070	Cu3247 ppm .0049 .0003 5.772	Zn2138 ppm 0002 .0004 177.4	Ni2316 ppm .0039 .0004 9.611	Tl1908 ppm 0128 .0130 101.5	Be3130 ppm .0025 .0001 3.301	Sb2068 ppm .0034 .0001 4.159
#1 #2	.0055	.0047	0006 .0001	.0042	0221 0036	.0026	.0033
Elem Unit Avge SDev %RSD	s ppm 0001 .0021	S_1820 ppm 15.53 .28 1.773	S_2203 ppm 0003 .0050 1712.	S_1960 ppm .0327 .0045 13.62	S_2068 ppm .0020 .0015 75.79	Ca3179 ppm 11.74 .21 1.788	Mg2790 ppm 10.58 .27 2.559
#1 #2	0015 .0014	15.72 15.34	0039 .0033	.0296 .0359	.0009	11.89 11.59	10.77 10.39
Elem Unit Avge SDev %RSD	s ppm .0001 .0000	Co2286 ppm .0002 .0004 212.1	Fe2714 ppm 0469 .0032 6.730	Al3082 ppm .4738 .0157 3.305	V_2924 ppm 0009 .0002 25.46	Mo2020 ppm .0023 .0005 23.02	Ti3349 ppm 0001 .0001 94.29
#1 #2	.0001	0001 .0005	0447 0491	.4848	0011 0008	.0020	0000 0002
Elem Unit Avge SDev %RSD	e .0001 .0005	B_2496 ppm 0374 .0004 .9698	K_7664 ppm 1.189 .010 .8106	Na3302 ppm 27.97 .99 3.542	Si2881 ppm .5834 .0102 1.748	Sn1899 ppm .0001 .0004 318.6	P_1782 ppm .0155 .0070 45.19
#1 #2	0002 .0004	0372 0377	1.196 1.182	28.67 27.27	.5906 .5762	0002 .0004	.0105

1A-IN
INORGANIC ANALYSIS DATA SHEET

RP-JM-SED-01	

Lab Name: Summit En	nvironmental Tech	Contract:	Jackpile	
Matrix: (soil/water)	soil	_ Lab Sample ID:	1107985-24	
Date Received:	04/26/2011	_		
% Solids:	85.42			

Concentration Units (mg/L or mg/kg dry weight): mg/kg

CAS No.	Analyte	Concentration	С	Q	M
7429-90-5	Aluminum	7230		D	Р
7440-36-0	Antimony	0.0937	U		Р
7440-38-2	Arsenic	6.57			P
7440-39-3	Barium	116			Р
7440-41-7	Beryllium	0.615	J		P
7440-43-9	Cadmium	0.0234	Ū		P
7440-70-2	Calcium	30400		D	P
7440-47-3	Chromium	9.33			P
7440-48-4	Cobalt	6.29			P
7440-50-8	Copper	6.18			P
7439-89-6	Iron	32600		D	P
7439-92-1	Lead	1.90	J		P
7439-95-4	Magnesium	4650			Р
7439-96-5	Manganese	490			P
7440-02-0	Nickel	10.6			Р
7440-09-7	Potassium	1950			P
7782-49-2	Selenium	30.0			P
7440-22-4	Silver	0.117	Ū		P
7440-23-5	Sodium	2330		D	P
7440-28-0	Thallium	0.187	Ū		P
7440-62-2	Vanadium	23.8			Р
7440-66-6	Zinc	34.1			P
7439-97-6	Mercury	0.0468	U		AV
		I			

Comments:						
	 	 		-		
	 	 	1*** #odes911	-	 	

Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-24s Run Time: 04/29/11 20:09:26

Comment: Mode: CC	6010/200	.7 CPI 440 . Factor:	0-013 lot	10B101	58.53	>	
Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.1598	.0780	03881	.1594	1.978	.1122	.5131
SDev	.0037	.0007	.0001	.0009	.007	.0032	.0035
%RSD	2.285	.8999	.2824	.5732	.3564	2.850	.6747
#1	.1624	.0785	0387	.1601	1.983	.1144	.5155
#2	.1573	.0775	0389	.1588	1.973	.1099	.5106
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	3420	.1056	.5825	.1803	0053	.0105	.0041
SDev	.0110	.0003	.0030	.0016	.0053	.0002	.0020
%RSD	3.219	.2336	.5103	.9167	99.74	1.525	49.88
#1	3498	.1057	.5846	.1814	0016	.0106	.0026
#2	3342	.1054	.5804	.1791	0090	.0104	.0055
Elem Units Avge SDev %RSD	Sb206A ppm 0752 .0007	S_1820 ppm 48.33 .21 .4387	S_2203 ppm .0325 .0014 4.355	S_1960 ppm .6067 .0039 .6349	S_2068 ppm0475 .0002 .4077	Ca3179 ppm H502.7 2.6	Mg2790 ppm 79.98 .31 .3838
#1	0747	48.48	.0336	.6039	0477	H504.5	80.19
#2	0757	48.18	.0315	.6094	0474	H500.8	79.76
Elem	Mn2576	Co2286	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	/ ppm	ppm	ppm	ppm
Avge	8.369	.1075	H459.5	H112.6	.4072	.0219	5.959
SDev	.027	.0006	1.7	.5	.0022	.0021	.024
%RSD	.3262	.5435	.3669	.4276	.5368	9.587	.4016
#1	8.389	.1079	H460.7	H112.9	.4087	.0234	5.976
#2	8.350	.1071	H458.3	H112.2	.4057		5.942
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0006	.8317	33.31	66.96	5.583	.0337	10.80
SDev	.0000	.0054	.18	.32	.018	.0017	.05
%RSD	4.705	.6478	.5410	.4798	.3304	4.937	.4875
#1	0006	.8355	33.44	67.19	5.597	.0325	10.84
#2	0006	.8279	33.18	66.73	5.570		10.76

Method: ZZJEM1_A Sample Name: 7985-24s x10
Run Time: 05/11/11 14:52:51
Comment: 6010/200.7 CPI 4400-013 lot 10B101
Mode: CONC Corr. Factor: 1 Operator: VVK

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0207	.0179	0006	.0193	.2231	.0107	.0537
SDev	.0022	.0014	.0002	.0001	.0010	.0076	.0013
%RSD	10.60	7.887	29.43	.3860	.4298	70.99	2.383
#1	.0223	.0189	0007	.0194	.2237	.0160	.0546
#2	.0192	.0169	0004	.0193	.2224	.0053	.0528
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	3547	.0171	.1005	.0254	0145	.0028	.0075
SDev	.0037	.0000	.0018	.0025	.0139	.0000	.0014
%RSD	1.033	.2629	1.816	9.704	96.06	.8461	18.65
#1	3521	.0171	.0992	.0272	0046	.0028	.0085
#2	3573	.0171	.1018	.0237	0243		.0065
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0048	3.250	.0103	1074	.0004	51.90	9.757
SDev	.0026	.007	.0017	.0032	.0013	.29	.036
%RSD	53.55	.2047	16.69	2.989	353.5	.5548	.3673
#1	0067	3.245	.0115	1051	0006	52.10	9.782
#2	0030	3.255		1096	.0013	51.69	9.732
Elem	Mn2576	Co2286	Fe2714	A13082	V_2924	Mo2020	Ti3349
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	1.020	.0141	55.73	12.36	.0479	.0016	.6860
SDev	.005	.0001	.24	.06	.0003	.0018	.0035
%RSD	.4770	.7068	.4384	.4471	.6829	118.5	.5078
#1	1.024	.0140	55.90	12.40	.0477	.0028	.6885
#2	1.017	.0142	55.55	12.32	.0482		.6835
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881 ppm .7131 .0007 .1036	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm		ppm	ppm
Avge	.0002	.0712	2.945	3.973		.0011	.9911
SDev	.0003	.0009	.012	.008		.0000	.0053
%RSD	143.3	1.212	.4011	.1987		.3673	.5368
#1	0000	.0718	2.937	3.968	.7136	.0011	.9873
#2	.0005	.0706	2.954	3.979	.7126		.9948

RP-JM-SED-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name:	Summit Environmental Tech	Contract:	Jackpile
Matrix: (soil	/water)soil	Lab Sample ID: _	1107985-25
Date Received	:04/26/2011		
% Solids:	76.14		
Concentration	. Units (mg/L or mg/kg dry weigl	h+), mg/kg	
Concentration	ource (mg/n or mg/kg ary werd)	$\frac{11C}{11}$: $\frac{111G}{K}$	

CAS No.	Analyte	Concentration	С	Q	Iv
7429-90-5	Aluminum	13500		D	P
7440-36-0	Antimony	0.105	U		P
7440-38-2	Arsenic	5.73			Р
7440-39-3	Barium	72.9			P
7440-41-7	Beryllium	0.729	J		P
7440-43-9	Cadmium	0.0263	U		P
7440-70-2	Calcium	18200		D	Р
7440-47-3	Chromium	11.7			P
7440-48-4	Cobalt	5.63			P
7440-50-8	Copper	9.81			P
7439-89-6	Iron	19800		D	P
7439-92-1	Lead	3.15	J		P
7439-95-4	Magnesium	6430			P
7439-96-5	Manganese	166			P
7440-02-0	Nickel	11.1			P
7440-09-7	Potassium	3410			P
7782-49-2	Selenium	23.6			P
7440-22-4	Silver	0.131	U		P
7440-23-5	Sodium	1520		D	P
7440-28-0	Thallium	0.210			P
7440-62-2	Vanadium	24.3	Ū		P
7440-66-6	Zinc	52.3	****		P
7439-97-6	Mercury	0.0525	U		AV

Com	ment	s:											
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Method: ZZJEM1_A Sample Name: 7985-25s

Run Time: 04/29/11 20:13:00

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

65	167	
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Operator: VVK

Elem Units Avge SDev %RSD	Pb2203 ppm .1933 .0000	Pb220A ppm .0342 .0020 5.747	Cd2265 ppm0203	Cr2677 ppm .1787 .0002 .1115	Ba4934 ppm 1.110 .003 .2765	As1890 ppm .0873 .0010 1.159	Se1960 ppm .2911 .0084 2.889
#1	.1933	.0329	0201	.1786	1.107	.0866	.2852
#2	.1933	.0356	0204	.1789	1.112	.0880	.2971
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	1670	.1494	.7963	.1693	0048	.0111	.0058
SDev	.0014	.0006	.0016	.0021	.0236	.0000	.0014
%RSD	.8474	.3773	.2004	1.231	493.3	.3656	23.44
#1 #2	1660 1680	.1490 .1498	.7974 .7951	.1678 .1708	0214 .0119	.0111	.0049
Elem Units Avge SDev %RSD	Sb206A ppm 0351 .0010 2.811	S_1820 ppm 22.42 .03 .1213	S_2203 ppm .0480 .0013 2.592	S_1960 ppm .3590 .0051 1.412	S_2068 ppm 0202 .0011 5.494	Ca3179 ppm H267.6 .6 .2322	Mg2790 ppm 97.89 .20
#1	0357	22.40	.0472	.3554	0210	H267.1	97.75
#2	0344	22.44	.0489	.3625	0195	H268.0	98.03
Elem Units Avge SDev %RSD	Mn2576 ppm 2.527 .008 .3006	Co2286 ppm .0857 .0003 .3125	Fe2714 ppm H248.6 .6 .2325	Al3082 ppm H190.6 .50 .2730	V_2924 ppm .3709 .0003 .0778	Mo2020 ppm .0143 .0004 2.603	Ti3349 ppm 1.053 .000
#1	2.522	.0856	H248.2	H190.3	.3707	.0146	1.053
#2	2.533	.0859	H249.0	H191.0		.0140	1.054
Elem	Ag3280	B_2496	K_7664	Na3302	/ Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0015	.5334	51.92	34.71	8.974	.0454	6.587
SDev	.0003	.0016	.07	.61	.019	.0005	.016
%RSD	21.54	.2995	.1276	1.743	.2173	1.056	.2470
#1	.0017	.5323	51.97	35.14	8.960	.0458	6.576
#2		.5345	51.87	34.28	8.988	.0451	6.599

Method: ZZJEM1_A Sample Name: 7985-25s x10 Run Time: 05/11/11 14:56:26

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0244	.0161	.0003	.0222	.1242	.0048	.0310
SDev	.0018	.0016	.0005	.0001	.0006	.0016	.0104
%RSD	7.434	10.19	161.7	.6756	.5146	33.33	33.50
#1	.0257	.0172	.0006	.0223	.1247	.0036	.0383
#2	.0231	.0149	0000	.0221	.1238	.0059	.0236
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	1922	.0210	.1752	.0258	0060	.0028	.0041
SDev	.0027	.0002	.0043	.0006	.0004	.0001	.0003
%RSD	1.423	1.071	2.474	2.354	6.428	1.786	7.142
#1 #2	1902 1941	.0211	.1783 .1722	.0262 .0254	0057 0063	.0028	.0039
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0006	1.617	.0143	0454	.0028	27.72	11.70
SDev	.0042	.006	.0017	.0054	.0029	.17	.01
%RSD	754.2	.3645	11.73	11.93	105.9	.6145	.0746
#1	0024	1.613	.0155	0416	.0007	27.84	11.71
#2	.0036	1.621		0492	.0048	27.60	11.69
Elem	Mn2576	Co2286 ppm .0102 .0004 4.359	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm		ppm	ppm	ppm	ppm	ppm
Avge	.3054		30.17	20.59	.0431	.0017	.1235
SDev	.0008		.09	.04	.0013	.0001	.0008
%RSD	.2440		.3074	.1895	3.038	3.449	.6068
#1	.3060	.0105	30.24	20.61	.0440	.0017	.1241
#2	.3049	.0099	30.10	20.56		.0018	.1230
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	7 ppm	ppm	ppm
Avge	.0012	.0421	4.968	2.322	1.182	.0020	.6654
SDev	.0001	.0011	.055	.008	5 .001	.0001	.0044
%RSD	5.155	2.542	1.098	.3404	.0804	5.910	.6639
#1 #2	.0012	.0428	5.007 4.930	2.328 2.317	1.181	.0020 .0021	.6685 .6623

110419

1A-IN INORGANIC ANALYSIS DATA SHEET

RP-JM-SW-01-110420

Lab Name:	Summit Envir	onmental Te	ch Contr	act:		Jackpi.	le
Matrix: (soil	./water)	water	Lab S	ample	e ID:	11079	85-26
Date Received							
		01/20/2011	· · · · · · · · · · · · · · · · · · ·				
% Solids:			Seattle				
Concentration	n Units (mg/	L or mg/kg	dry weight): _m	g/L	_		
	CAS No.	Analyte	Concentration	С		Q	М
	7429-90-5	Aluminum	0.0105	U			P
	7440-36-0	Antimony	0.00237	U			Р
	7440-38-2	Arsenic	0.00206	U			Р
	7440-39-3	Barium	0.0370	J			Р
	7440-41-7	Beryllium	0.00072	Ū			Р
	7440-43-9	Cadmium	0.00082	U			Р
	7440-70-2	Calcium	200		D		Р
	7440-47-3	Chromium	0.00109	U			P
	7440-48-4	Cobalt	0.00051	U			P
	7440-50-8	Copper	0.00183	Ū			Р
	7439-89-6	Iron	0.0106	U			Р
	7439-92-1	Lead	0.00210	U			Р
	7439-95-4	Magnesium	150		D		Р
	7439-96-5	Manganese	0.0348				Р
	7440-02-0	Nickel	0.00116	U		*******	Р
	7440-09-7	Potassium	15.3				Р
	7782-49-2	Selenium	0.0341				Р
	7440-22-4	Silver	0.00039	Ū			P
	7440-23-5	Sodium	295		D		P
	7440-28-0	Thallium	0.0071	J			Р
	7440-62-2	Vanadium	0.00174	Ü			Р
	7440-66-6	Zinc	0.0025	J			Р
	7439-97-6	Mercury	0.00006	Ū			AV

Com	ments:						
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Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-26

Run Time: 04/29/11 19:30:09

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0015 .0019 131.2	Pb220A ppm 0023 .0004 17.03	Cd2265 ppm 0001 .0000 35.10	Cr2677 ppm 0005 .0000 3.126	Ba4934 ppm .0370 .0002 .4360	As1890 ppm .0000 .0026 9158.	Se1960 ppm .0106 .0054 51.40
#1 #2	.0028	0021 0026	0001 0000	0005 0005	.0369 .0371	0018 .0019	.0144
Elem Units Avge SDev %RSD	Se196A ppm .0025 .0014 54.00	Cu3247 ppm 0048 .0010 19.78	Zn2138 ppm .0025 .0001 4.007	Ni2316 ppm .0002 .0006 342.1	Tl1908 ppm .0071 .0013 18.93	Be3130 ppm 0001 .0000	Sb2068 ppm 0038 .0064 167.8
#1 #2	.0015	0055 0041	.0025	.0006 0003	.0061	0001 0001	0083 .0007
Elem Units Avge SDev %RSD	Sb206A ppm .0012 .0036 289.2	S_1820 ppm H399.9 2.4 .6018	S_2203 ppm 0008 .0009 117.7	S_1960 ppm .0341 .0015 4.379	S_2068 ppm .0005 .0045 833.8	Ca3179 ppm H202.0 1.5 .7284	Mg2790 ppm H142.7 1.0
#1 #2	0013 .0037	H398.2 H401.6	0001 0014	.0330	0026 .0037	H200.9 H203.0	H142.0 H143.4
Elem Units Avge SDev %RSD	Mn2576 ppm .0348 .0001 .2446	Co2286 ppm 0002 .0004 192.9	Fe2714 ppm 0504 .0007 1.479	Al3082 ppm 0362 .0008 2.128	V_2924 ppm 0152 .0002 1.519	Mo2020 ppm .0032 .0015 47.29	Ti3349 ppm0023 .0001 5.262
#1 #2	.0347	.0001 0005	0510 0499	0368 0357	0150 0153	.0022	0022 0024
Elem Units Avge SDev %RSD	Ag3280 ppm 0005 .0006 122.6	B_2496 ppm .1159 .0010 .8340	K_7664 ppm 15.33 .12 .7589	Na3302 ppm H427.8 3.6 .8515	Si2881 ppm 8.107 .049 .5985	Sn1899 ppm .0040 .0004 11.23	P_1782 ppm .0656 .0005 .7452
#1 #2	0001 0008	.1152 .1166	15.25 15.41	H425.3 H430.4	8.073 8.141	.0037	.0659 .0652

Method: ZZJEM1_A Sample Name: 7985-26 x10 Run Time: 05/11/11 13:14:02 Operator: VVK

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm 0017 .0100 585.9	Pb220A ppm .0030 .0056 190.5	Cd2265 ppm .0001 .0004 271.3	Cr2677 ppm .0013 .0001 7.859	Ba4934 ppm .0044 .0001 1.589	As1890 ppm 0074 .0064 87.15	Se1960 ppm 0032 .0078 242.4
#1 #2	.0054 0088	.0069 0010	.0004 0001	.0012	.0043	0119 0028	.0023 0087
Elem Units Avge SDev %RSD	Se196A ppm .0033 .0021 63.03	Cu3247 ppm .0055 .0001 .8623	Zn2138 ppm 0002 .0003 169.7	Ni2316 ppm .0020 .0022 114.2	Tl1908 ppm 0155 .0065 42.13	Be3130 ppm .0024 .0002 6.884	Sb2068 ppm .0030 .0018 61.28
#1 #2	.0018	.0054 .0055	.0000	.0036	0201 0109	.0025	.0017
Elem Units Avge SDev %RSD	Sb206A ppm .0001 .0017 1556.	S_1820 ppm 20.95 .02 .0771	S_2203 ppm .0018 .0071 401.1	S_1960 ppm .0283 .0012 4.163	S_2068 ppm .0021 .0017 84.56	Ca3179 ppm 19.99 .02 .0933	Mg2790 ppm 15.01 .00
#1 #2	0011 .0013	20.96 20.94	.0068 0033	.0291 .0275	.0008	19.98 20.01	15.02 15.01
Elem Units Avge SDev %RSD	Mn2576 ppm .0051 .0000 .0334	Co2286 ppm .0007 .0003 47.18	Fe2714 ppm 0429 .0123 28.54	Al3082 ppm .4709 .0151 3.200	V_2924 ppm 0006 .0000 .4890	Mo2020 ppm .0023 .0006 26.94	Ti3349 ppm .0002 .0000
#1 #2	.0051 .0051	.0009	0516 0343	.4815 .4602	0006 0006	.0018	.0002
Elem Units Avge SDev %RSD	Ag3280 ppm .0003 .0001 54.98	B_2496 ppm 0394 .0005 1.341	K_7664 ppm 1.213 .016 1.327	Na3302 ppm 29.46 .08 .2732	Si2881 ppm 1.008 .001 .1264	Sn1899 ppm 0000 .0005 5380.	P_1782 ppm .0215 .0051 23.83
#1 #2	.0004	0391 0398	1.225 1.202	29.40 29.51	1.009	.0004	.0179 .0252

110419

1A-IN INORGANIC ANALYSIS DATA SHEET

RP-JM-SW-110420

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ΑV

Lab Name:	Summit Envir	conmental Te	ch Contr	act:	Jackpi	.le
Matrix: (soi	/water)	water	Lab S	ample	e ID:11079	985-27
Date Received	d:	04/26/2011				
% Solids:						
Concentration	<u> </u>	T	dry weight): _m	1		
	CAS No.	Analyte	Concentration	С	Q	M
	7429-90-5	Aluminum	0.0105	Ū		P
	7440-36-0	Antimony	0.0048	J		P
	7440-38-2	Arsenic	0.00206	U		P
	7440-39-3	Barium	0.0472			P
	7440-41-7	Beryllium	0.00072	Ų		P
	7440-43-9	Cadmium	0.00082	U		P
	7440-70-2	Calcium	181		D	P
	7440-47-3	Chromium	0.00109	U		P
	7440-48-4	Cobalt	0.00051	Ū		P
	7440-50-8	Copper	0.00183	Ū		P
	7439-89-6	Iron	0.0106	Ū		P
	7439-92-1	Lead	0.00210	Ū		P
	7439-95-4	Magnesium	96.7		D	P
	7439-96-5	Manganese	0.0374			P

0.0019

0.0276

0.0012

0.00529

0.00174

0.0034

0.00006

127

12.3

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7440-02-0 Nickel

7440-22-4 Silver

7440-23-5 Sodium

7440-66-6 Zinc

7440-28-0 Thallium

7440-62-2 Vanadium

7439-97-6 Mercury

7440-09-7 Potassium

7782-49-2 | Selenium

Com	ments:					
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Operator: VVK

Method: ZZJEM1_A Sample Name: 7985-27 Run Time: 04/29/11 19:33:43

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0007	0041	.0001	.0003	.0472	0005	.0043
SDev	.0066	.0016	.0001	.0003	.0003	.0037	.0017
%RSD	1019.	39.52	81.80	103.4	.6843	799.5	39.90
#1 #2	.0053 0040	0029 0052	.0000	.0001 .0006	.0470 .0474	.0022 0031	.0055
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0015	0032	.0034	.0019	.0049	0000	0027
SDev	.0036	.0001	.0002	.0018	.0116	.0000	.0048
%RSD	249.6	4.351	5.301	94.28	237.8	112.6	175.1
#1 #2	.0011	0033 0031	.0033	.0006 .0032	.0131	0001 0000	.0006 0061
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0070	H231.8	0022	.0276	.0048	H179.6	90.10
SDev	.0086	2.1	.0032	.0030	.0041	1.4	.73
%RSD	122.2	.9003	149.2	10.83	86.48	.7785	.8062
#1	.0010	H230.3	.0001	.0297	.0018	H178.6	89.58
#2		H233.3	0045	.0255	.0077	H180.6	90.61
Elem	Mn2576	Co2286 ppm .0005 .0001 30.77	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm		ppm	ppm	ppm	ppm	ppm
Avge	.0374		0599	0366	0061	.0034	0020
SDev	.0004		.0027	.0099	.0005	.0011	.0001
%RSD	1.189		4.567	27.02	8.217	33.07	2.961
#1 #2	.0371	.0006	0580 0619	0436 0296	0064 0057	.0026 .0042	0021 0020
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0012	.0558	12.25	H174.8	10.67	.0063	.0810
SDev	.0007	.0010	.07	1.1	.10	.0001	.0061
%RSD	56.88	1.713	.6094	.6131	.8943	1.293	7.585
#1	.0017	.0565	12.20	H174.1	10.60	.0062	.0767
#2		.0552	12.31	H175.6	10.73	.0063	.0854

Method: ZZJEM1_A Sample Name: 7985-27 x10 Operator: VVK Run Time: 05/11/11 13:24:45 Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

Elem Units Avge SDev %RSD	Pb2203 ppm .0002 .0026 1144.	Pb220A ppm 0028 .0041 144.5	Cd2265 ppm .0003 .0002 64.16	Cr2677 ppm .0008 .0004 51.46	Ba4934 ppm .0059 .0000	As1890 ppm .0023 .0003 13.34	Se1960 ppm .0102 .0034 33.83
#1 #2	0016 .0020	0057 .0001	.0002	.0011	.0059	.0025 .0021	.0077 .0126
Elem	Se196A	Cu3247	Zn2138	Ni2316	Tl1908 ppm0016 .0074 452.5	Be3130	Sb2068
Units	ppm	ppm	ppm	ppm		ppm	ppm
Avge	.0133	.0055	.0004	.0031		.0023	.0132
SDev	.0041	.0001	.0001	.0003		.0001	.0054
%RSD	30.53	.8623	12.39	8.730		4.596	40.78
#1 #2	.0104	.0055	.0005	.0033	0069 .0036	.0023	.0170
Elem Units Avge SDev %RSD	Sb206A ppm .0041 .0021 50.91	S_1820 ppm 12.39 .07 .5599	S_2203 ppm 0015 .0035 240.0	S_1960 ppm .0394 .0038 9.725	S_2068 ppm .0081 .0032 39.18	Ca3179 ppm 18.10 .16 .8892	Mg2790 ppm 9.671 .087
#1	.0056	12.44	0040	.0367	.0104	18.21	9.732
#2	.0026	12.34	.0010	.0422	.0059	17.98	9.609
Elem	Mn2576	Co2286 ppm0000 .0004 59370.	Fe2714	Al3082	V_2924	Mo2020	Ti3349
Units	ppm		ppm	ppm	ppm	ppm	ppm
Avge	.0055		0219	.4189	0003	.0023	.0010
SDev	.0001		.0109	.0017	.0000	.0011	.0001
%RSD	1.069		49.85	.4151	.9869	47.14	5.439
#1	.0056	0003	0142	.4176	0003	.0030	.0010
#2	.0055	.0003	0297	.4201	0003	.0015	
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.0012	0400	.9328	12.68	1.329	.0003	.0169
SDev	.0003	.0004	.0078	.08	.006	.0003	.0019
%RSD	20.46	.9461	.8386	.6496	.4890	100.8	11.50
#1 #2	.0014	0397 0402	.9384 .9273	12.62 12.74	1.334 1.325	.0001	.0156 .0183

RP-SED-01-110419

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name:	Summit En	vironmental Tech	Contract:	Jackpile	
Matrix: (soi	l/water) _	soil	Lab Sample ID: _	1107985-28	
Date Receive	d:	04/26/2011			
% Solids:		78.35			

Concentration Units (mg/L or mg/kg dry weight): mg/kg

CAS No.	Analyte	Concentration	С	Q	М
7429-90-5	Aluminum	10100		D	P
7440-36-0	Antimony	0.102	U		P
7440-38-2	Arsenic	4.56			Р
7440-39-3	Barium	76.2			P
7440-41-7	Beryllium	0.574	J		P
7440-43-9	Cadmium	0.0255	U		P
7440-70-2	Calcium	26000		D	Р
7440-47-3	Chromium	8.09			Р
7440-48-4	Cobalt	4.00			Р
7440-50-8	Copper	7.26			Р
7439-89-6	Iron	15500		D	P
7439-92-1	Lead	3.32	J		Р
7439-95-4	Magnesium	8410		D	P
7439-96-5	Manganese	150			Р
7440-02-0	Nickel	7.54			Р
7440-09-7	Potassium	4060			P
7782-49-2	Selenium	18.4			P
7440-22-4	Silver	0.128	Ū		P
7440-23-5	Sodium	4830		D	P
7440-28-0	Thallium	0.747	J		P
7440-62-2	Vanadium	18.6			P
7440-66-6	Zinc	32.7			P
7439-97-6	Mercury	0.0511	Ū		AV

Comme	nts:								
	*		 	 10/3		 		- -	
-				 13.774		 .			
			 	 	11.00		***	,	

Method: ZZJEM1_A Sample Name: 7985-28s

Run Time: 04/29/11 20:16:34

Comment: 6010/200.7 CPI 4400-013 lot 10B101 Mode: CONC Corr. Factor: 1

63.82

Operator: VVK

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.1672	.0423	0165	.1267	1.194	.0715	.2247
SDev	.0050	.0003	.0001	.0004	.001	.0013	.0044
%RSD	2.961	.6799	.8794	.3154	.0496	1.810	1.968
#1	.1707	.0425	0164	.1264	1.193	.0706	.2278
#2	.1637	.0421	0166	.1270	1.194	.0725	.2215
Elem Units Avge SDev %RSD	Se196A ppm 1427 .0166 11.61	Cu3247 ppm .1138 .0013	Zn2138 ppm .5117 .0001 .0295	Ni2316 ppm .1182 .0004 .3162	Tl1908 ppm .0117 .0088 75.42	Be3130 ppm .0090 .0000	Sb2068 ppm .0011 .0020 179.8
#1 #2	1544 1310	.1130	.5116 .5118	.1179 .1185	.0179	.0090	.0025
Elem	Sb206A	S_1820	S_2203	S_1960	S_2068	Ca3179	Mg2790
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0330	H187.8	.0520	.2882	0205	H404.7	H108.5
SDev	.0015	.1	.0019	.0097	.0016	.1	.1
%RSD	4.405	.0324	3.574	3.365	7.975	.0210	.0866
#1	0319	H187.9	.0533	.2814	0193	H404.6	H108.4
#2	0340	H187.8	.0506	.2951	0216	H404.8	H108.5
Elem Units Avge SDev %RSD	Mn2576 ppm 2.357 .001 .0584	Co2286 ppm .0626 .0001 .1589	Fe2714 ppm H203.1 .1	Al3082 ppm H147.7 .0 .0153	V_2924 ppm .2913 .0004 .1498	Mo2020 ppm .0121 .0013 11.12	Ti3349 ppm .9901 .0006 .0573
#1	2.356	.0625	H203.0	H147.7	.2910	.0111	.9897
#2	2.358	.0626	H203.2	H147.6	.2916		.9905
Elem Units Avge SDev %RSD	Ag3280 ppm 0015 .0002 13.82	B_2496 ppm .4315 .0027 .6308	K_7664 ppm 63.67 .03	Na3302 ppm H143.5 .2 .1567	Si2881 ppm 5.166 .001 .0285	Sn1899 ppm .0378 .0023 5.983	P_1782 ppm 6.051 .011 .1746
#1	0016	.4335	63.65	H143.7	5.165	.0362	6.058
#2	0013	.4296	63.70	H143.4	5.167		6.043

Method: ZZJEM1_A Sample Name: 7985-28s Run Time: 05/11/11 15:00:00 x10

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Elem Units Avge SDev %RSD	Pb2203 ppm .0195 .0081 41.54	Pb220A ppm .0152 .0019 12.70	Cd2265 ppm .0003 .0003 126.0	Cr2677 ppm .0158 .0000 .0006	Ba4934 ppm .1307 .0004 .2934	As1890 ppm .0006 .0059 1071.	Se1960 ppm .0272 .0078 28.65
#1 #2	.0252 .0137	.0166 .0139	.0000	.0158 .0158	.1310	.0048	.0327 .0217
Elem Units Avge SDev %RSD	Se196A ppm 1554 .0044 2.862	Cu3247 ppm .0174 .0004 2.585	Zn2138 ppm .0882 .0015 1.707	Ni2316 ppm .0194 .0024 12.29	Tl1908 ppm 0281 .0301 107.1	Be3130 ppm .0025 .0001 3.427	Sb2068 ppm .0037 .0039 103.7
#1 #2	1523 1586	.0171	.0871 .0892	.0211 .0177	0068 0495	.0024	.0065
Elem Units Avge SDev %RSD	Sb206A ppm 0035 .0041 118.5	S_1820 ppm 12.11 .06 .4621	S_2203 ppm .0131 .0039 29.80	S_1960 ppm 0354 .0032 9.176	S_2068 ppm 0001 .0016 3068.	Ca3179 ppm 40.73 .12 .2848	Mg2790 ppm 7 13.17 .03 .2481
#1 #2	0064 0006	12.15 12.07	.0158	0331 0376	0012 .0011	40.81 40.64	13.20 13.15
Elem Units Avge SDev %RSD	Mn2576 ppm .2807 .0011 .4062	Co2286 ppm .0085 .0004 5.260	Fe2714 ppm 24.31 .06 .2597	Al3082 ppm 15.88 .04 .2671	V_2924 ppm .0337 .0000 .0010	Mo2020 ppm .0004 .0005 113.1	Ti3349 ppm .1124 .0000
#1 #2	.2815 .2799	.0082	24.36 24.27	15.91 15.85	.0337	.0008	.1124
Elem Units Avge SDev %RSD	Ag3280 ppm .0000 .0000	B_2496 ppm .0199 .0016 8.208	K_7664 ppm 5.051 .005	Na3302 ppm 7.570 .010	Si2881 ppm .6577 .0016 .2407	Sn1899 ppm .0012 .0003 28.22	P_1782 ppm .5534 .0073 1.313
#1 #2	.0000	.0211	5.047 5.054	7.562 7.577	.6588 .6566	.0015	.5585 .5482

RP-SED-02-110420

1A-IN INORGANIC ANALYSIS DATA SHEET

Lab Name: Summit E	nvironmental Tech	Contract:	Jackpile	
Matrix: (soil/water)	soil	_ Lab Sample ID: _	1107985-29	
Date Received:	04/26/2011			
% Solids:	77.62	_		
Concentration Units	(mg/L or mg/kg dry we	ight): <u>mg/kg</u>		

CAS No.	Analyte	Concentration	С	Q	М
7429-90-5	Aluminum	13900		D	P
7440-36-0	Antimony	0.103	U		P
7440-38-2	Arsenic	4.52			P
7440-39-3	Barium	157			Р
7440-41-7	Beryllium	0.728	J		P
7440-43-9	Cadmium	0.0258	Ū		P
7440-70-2	Calcium	31700		D	P
7440-47-3	Chromium	9.58			P
7440-48-4	Cobalt	4.88			P
7440-50-8	Copper	8.87			Р
7439-89-6	Iron	16900		D	P
7439-92-1	Lead	2.59	J		P
7439-95-4	Magnesium	7920		D	P
7439-96-5	Manganese	258			P
7440-02-0	Nickel	8.95			P
7440-09-7	Potassium	4250			P
7782-49-2	Selenium	19.9			P
7440-22-4	Silver	0.129	Ū		P
7440-23-5	Sodium	1850		D	P
7440-28-0	Thallium	0.206	Ū		P
7440-62-2	Vanadium	22.2			P
7440-66-6	Zinc	38.8			P
7439-97-6	Mercury	0.0515	Ū		AV

Com	ments	:									
			301	-	 	 	-	-		 	
			 		 	 			_	-	

Method: ZZJEM1_A Sample Name: 7985-29s

Run Time: 04/29/11 20:20:09

Comment: 6010/200.7 CPI 4400-013 lot 10B101

Mode: CONC Corr. Factor: 1

64.42

Operator: VVK

Elem	Pb2203	Pb220A	Cd2265	Cr2677	Ba4934	As1890	Se1960
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	.1910	.0161	0176	.1487	2.441	.0701	.2485
SDev	.0007	.0014	.0001	.0006	.006	.0002	.0005
%RSD	.3751	8.353	.7311	.3804	.2337	.3463	.1947
#1	.1905	.0152	0175	.1491	2.445	.0703	.2482
#2	.1915	.0171	0176	.1483		.0699	.2489
Elem Units Avge SDev %RSD	Se196A ppm 1531 .0001	Cu3247 ppm .1377 .0004 .2814	Zn2138 ppm .6029 .0005 .0852	Ni2316 ppm .1389 .0012 .8845	Tl1908 ppm .0016 .0139 863.2	Be3130 ppm .0113 .0001 .4818	Sb2068 ppm0053 .0056 105.5
#1 #2	1531 1532	.1380 .1375	.6032 .6025	.1398	0082 .0114	.0113	0093 0013
Elem Units Avge SDev %RSD	Sb206A ppm 0315 .0003	S_1820 ppm 23.58 .05 .2120	S_2203 ppm .0402 .0012 2.981	S_1960 ppm .3087 .0007 .2149	S_2068 ppm 0216 .0021 9.585	Ca3179 ppm H482. 1.3 .2715	Mg2790 ppm H101.2 .2
#1	0318	23.61	.0394	.3092	0231	H483.9	H101.3
#2	0313	23.54		.3082	0202	H482.0	H101.0
Elem Units Avge SDev %RSD	Mn2576 ppm 4.004 .009 .2182	Co2286 ppm .0758 .0006 .8130	Fe2714 ppm H217.2 .5 .2464	Al3082 ppm H199.9 .6	V_2924 ppm .3442 .0009 .2656	Mo2020 ppm .0087 .0006 6.350	Ti3349 ppm .8661 .0023 .2656
#1	4.010	.0762	H217.5	H200.3	.3448	.0091	.8678
#2	3.997	.0754	H216.8	H199.4		.0083	.8645
Elem	Ag3280	B_2496	K_7664	Na3302	Si2881	Sn1899	P_1782
Units	ppm	ppm	ppm	ppm	ppm	ppm	ppm
Avge	0018	.4610	66.05	44.32	5.283	.0467	6.668
SDev	.0004	.0032	.26	.09	.007	.0005	.023
%RSD	20.94	.6996	.3971	.2029	.1260	1.145	.3523
#1	0015	.4633	66.24	44.26	5.288	.0471	6.651
#2	0021	.4587	65.87	44.39	5.278		6.684